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SPECIFICATION

TRANSITION METAL COMPLEX LIGAND AND OLEFIN POLYMERIZATION
CATALYST CONTAINING TRANSITION METAL COMPLEX

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Technical Field

The invention relates to a transition metal complex, a ligand and an olefin polymerization catalyst, and a production method of an olefin polymer.

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Background Art

Conventionally, it has been reported to use a reaction product of an organic compound having two hydroxyl groups and phosphine (for example,

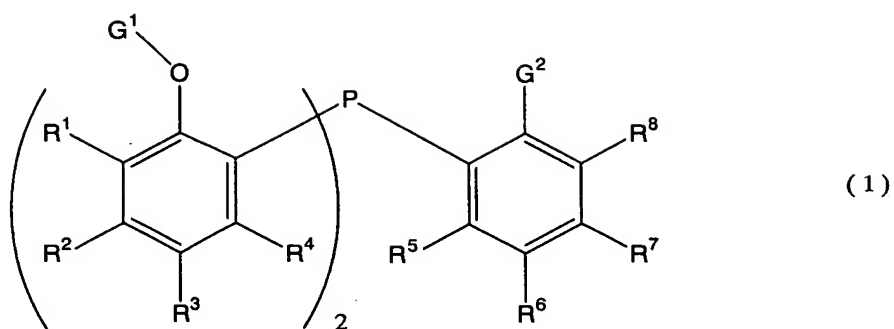
15 2,2'-(phenylphosphide)bis(6-tert-butyl-4-methylphenoxide)(tetrahydrofuran)titanium dichloride) for production of olefin polymers (for example Japanese Patent Application Laid-Open (JP-A) No. 10-218922).

20 Disclosure of the Invention

The transition metal complex having the ligand of the invention is useful as a component of an olefin polymerization catalyst. The catalyst has a good polymerization activity and can be used for production of high molecular weight olefin
25 polymers.

The present invention provides:

1. a phosphine compound of formula (1):



wherein $R^1, R^2, R^3, R^4, R^6, R^7$ and R^8 are the same or different, and independently represent;

a hydrogen atom,

5 a halogen atom,

a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

10 a substituted or unsubstituted aryl group having 6 to 20 carbon atoms,

a silyl group substituted with a substituted or unsubstituted hydrocarbon having 1 to 20 carbon atom(s),

15 a substituted or unsubstituted alkoxy group having 1 to 10 carbon atom(s),

a substituted or unsubstituted aralkyloxy group having 7 to 20 carbon atoms,

a substituted or unsubstituted aryloxy group having 6 to 20 carbon atoms, or

20 an amino group disubstituted with hydrocarbons having 1 to 20 carbon atom(s);

R^5 represents,

a hydrogen atom,

a fluorine atom,

a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

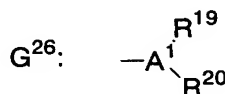
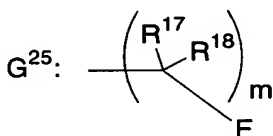
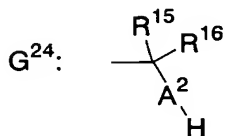
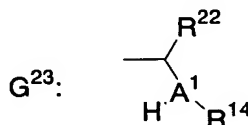
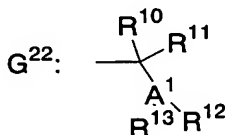
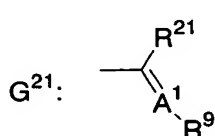
a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

5 a substituted or unsubstituted aryl group having 6 to 20 carbon atoms, or

a silyl group substituted with a substituted or unsubstituted hydrocarbon having 1 to 20 carbon atom(s);

10 G^1 represents a hydrogen atom or a protective group of hydroxyl group;

G^2 represents any one of G^{21} to G^{26} below;



15 wherein A^1 represents an element of Group 15 of the periodic table, and A^2 represents an element of Group 16 of the periodic table, wherein A^1 in G^{21} represents a nitrogen atom;

R^9 and R^{14} each represents

a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

20 a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

a substituted or unsubstituted aryl group having 6 to 20 carbon atoms, or

a group of formula:

$R^{90}-N-R^{91}$

wherein R^{90} and R^{91} are the same or different, and represent a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

5 a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

a substituted or unsubstituted aryl group having 6 to 20 carbon atoms, or

a cyclic structure by being linked together;

10 R^{12} , R^{13} , R^{19} and R^{20} each independently represents

a substituted or unsubstituted alkyl group 1 to 10,

a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

15 a substituted or unsubstituted aryl group having 6 to 20 carbon atoms, or

R^{12} and R^{13} , and R^{19} and R^{20} , each independently, are linked together and represent cyclic structure;

R^{10} , R^{11} , R^{15} , R^{16} , R^{21} and R^{22} each independently represents A hydrogen atom,

20 a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms, or

25 a substituted or unsubstituted aryl group having 6 to 20 carbon atoms,

R^{17} and R^{18} are the same or different, and represent,

a hydrogen atom,

a halogen atom,

a substituted or unsubstituted alkyl group having 1 to

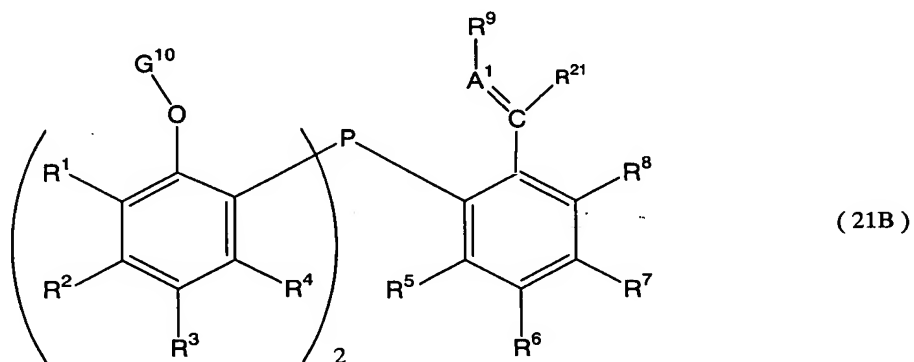
10 carbon atom(s),

a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms, or

a substituted or unsubstituted aryl group having 6 to 20 carbon atom(s); and

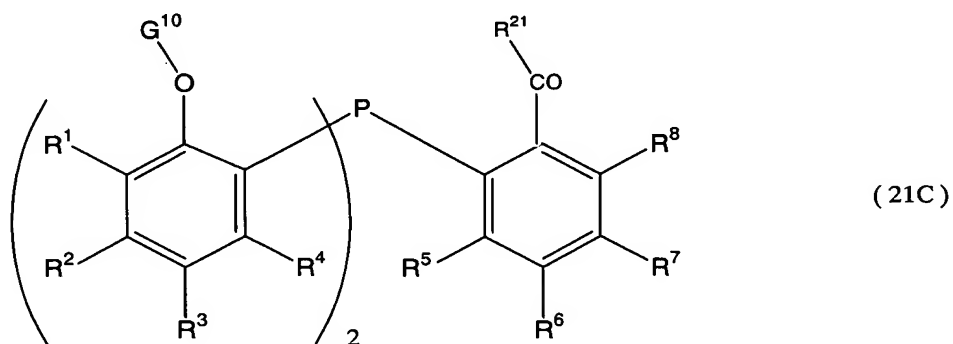
m represents an integer of 0 or 1;

2. a production method of a phosphine compound of formula (21B):



10 wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{21} and G^{10} are as defined below, which comprises reacting

a phosphine carbonyl compound of formula (21C):



15 wherein G^{10} represents a hydrogen atom, or a protective group of a hydroxyl group selected from an alkyl group having a secondary or tertiary carbon atom linked to an oxygen atom of phenol, or a C1 to C2 alkyl group substituted with a substituted

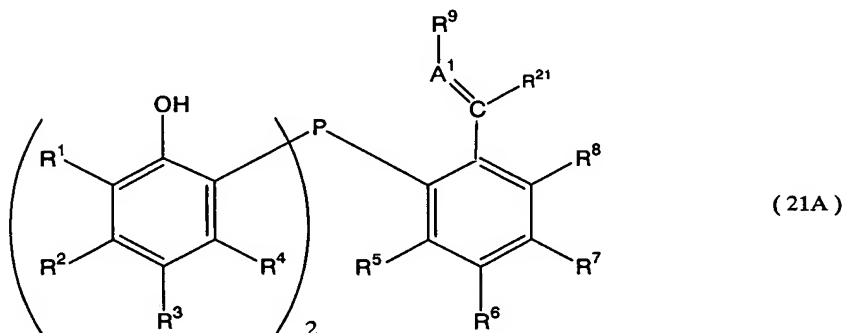
or unsubstituted alkoxy group, and

$R^1, R^2, R^3, R^4, R^5, R^6, R^7, R^8$ and R^{21} are the same as defined above, with an organic compound of formula (21F):



5 wherein R^9 is as defined above;

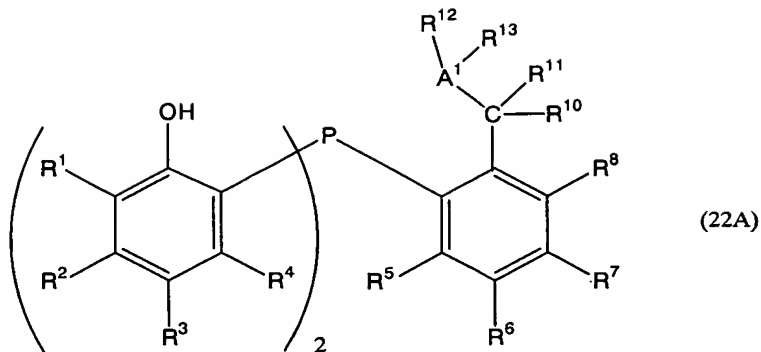
3. a production method of a phosphine compound of formula 21A:



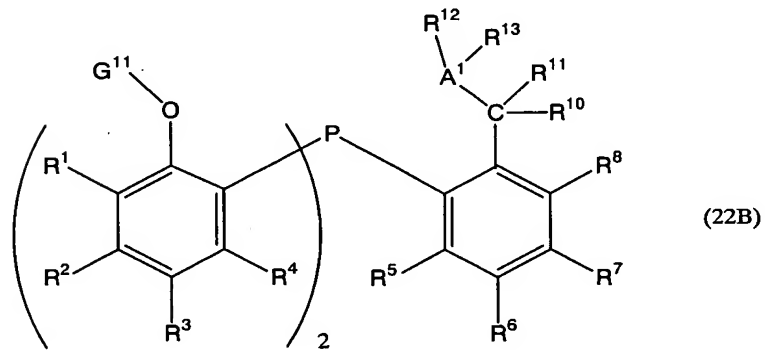
10 wherein $R^1, R^2, R^3, R^4, R^5, R^6, R^7, R^8$ and R^9 are as described above, and A^1 represents a nitrogen atom, which comprises

reacting the phosphine compound of formula 21B above, wherein G^{10} is a protective group of hydroxyl group selected from an alkyl group having a secondary or tertiary carbon atom linked to an oxygen atom of phenol or a C1 or C2 alkyl group substituted with a substituted or unsubstituted alkoxy group, with an acid;

4. a production method of the phosphine compound of formula (22A):



wherein $R^1, R^2, R^3, R^4, R^5, R^6, R^7, R^8, R^{10}, R^{11}, R^{12}, R^{13}$ and A^1 are as described above, which comprises reacting a phosphine compound of formula (22B):

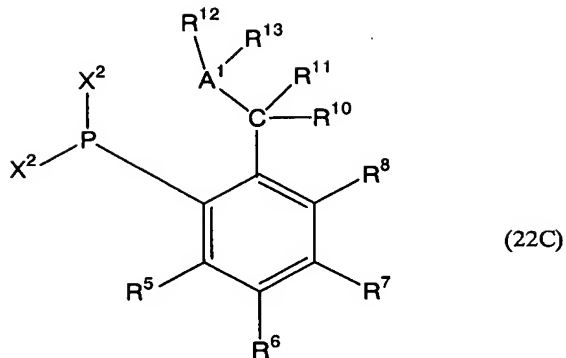


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wherein $R^1, R^2, R^3, R^4, R^5, R^6, R^7, R^8, R^{10}, R^{11}, R^{12}, R^{13}$ and A^1 are the same as described above, and G^{11} represents a protective group of hydroxyl group selected from an alkyl group having a secondary or tertiary carbon atom linked to an oxygen atom of phenol, or C1 to C2 alkyl groups substituted with a substituted or unsubstituted alkoxy group, with an acid;

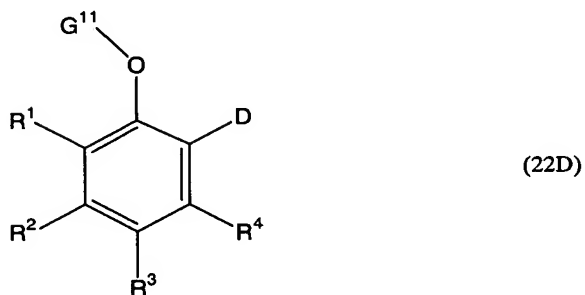
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5. a production method of a phosphine compound of formula (22B) above, which comprises reacting a phosphine dihalide compound of formula (22C):



wherein A, R⁵, R⁶, R⁷, R⁸, R¹¹, R¹², R¹³ and A¹ are as defined above, and X² represents a halogen atom,

with a metal aryl compound of formula (22D);

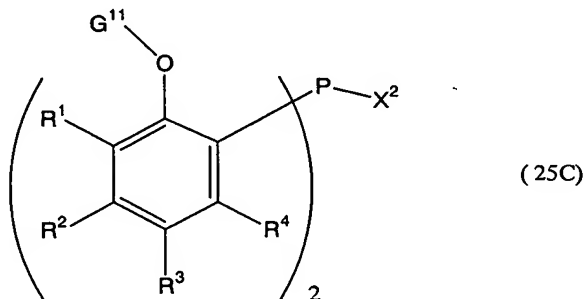


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wherein R¹, R², R³, R⁴ and G¹¹ are the same as above, and D represents an alkali metal or J-X³, where J represents an alkaline earth metal and X³ represents a halogen atom;

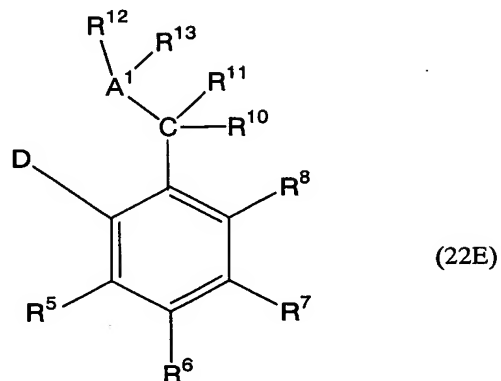
6. a production method of a phosphine compound of formula
10 (22B) above which comprises reacting

a phosphine halide compound of formula (25C):



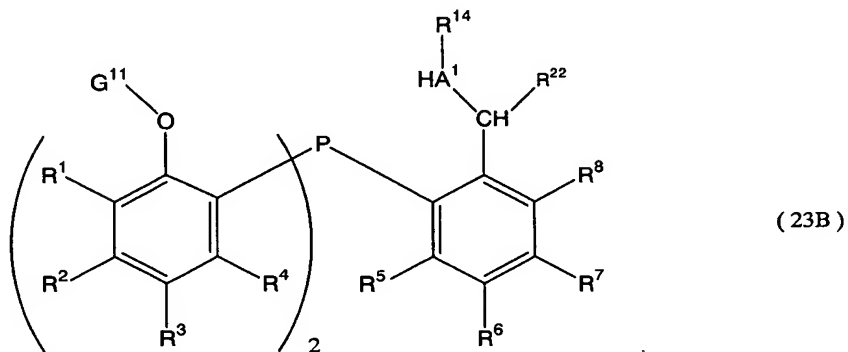
wherein R¹, R², R³, R⁴ and G¹¹ are as described above, and

X^2 represents a halogen atom, with a metal aryl compound of formula (22E):

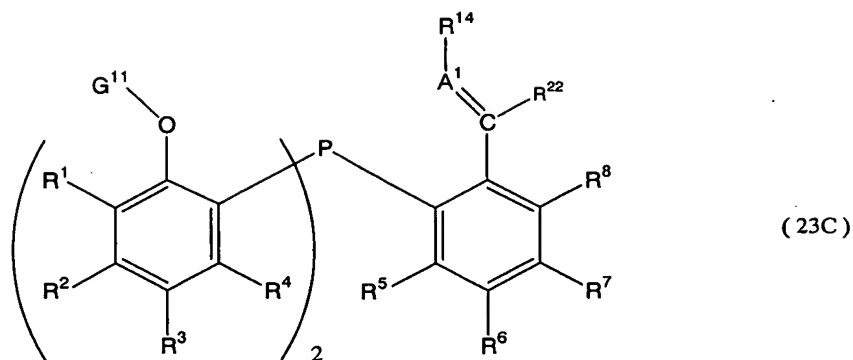


wherein A, R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} , R^{12} , R^{13} , A^1 and D are as described above;

7. a production method of a phosphine compound of formula (23B):

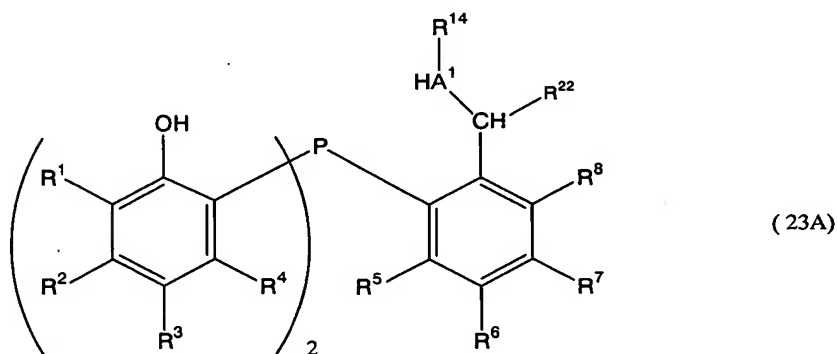


wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{14} , A^1 and R^{22} are as described above, which comprises reacting a phosphine compound of formula (23C):



wherein A^1 , R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{14} , G^{11} and R^{22} are as described above, with a metal hydride compound;

8. a production method of a phosphine compound of formula
5 (23A):

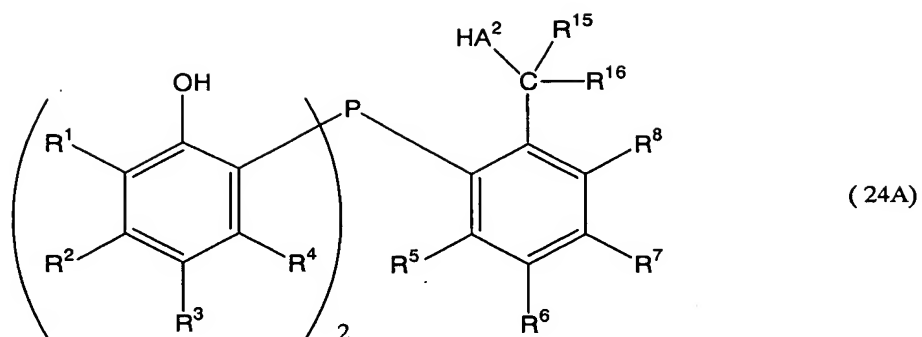


wherein , A^1 , R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{14} , A^1 and R^{22} are as described above,

which comprises reacting

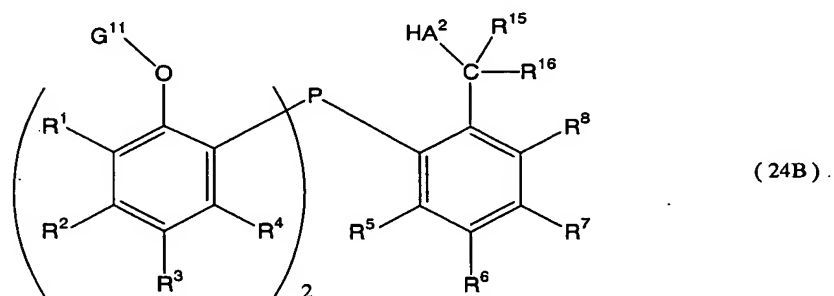
10 a phosphine compound of formula (23B) with an acid;

9. a production method of a phosphine compound of formula
(24A):



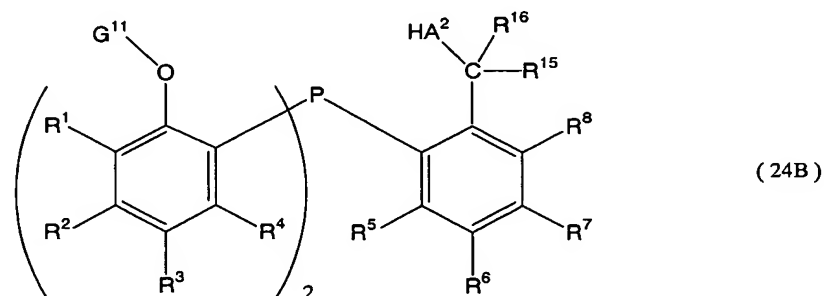
wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{15} , R^{16} and A^2 are as described above,

5 which comprises reacting a phosphine compound of formula (24B) :



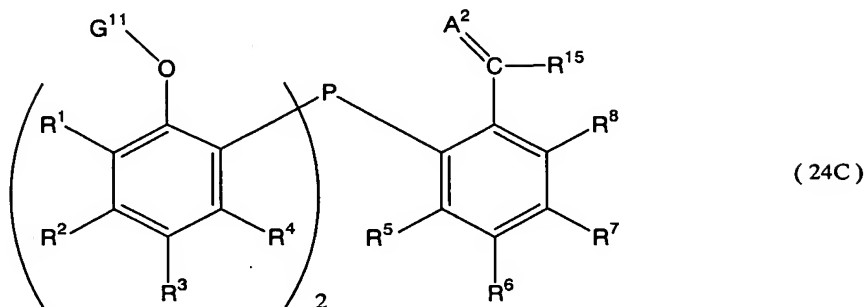
wherein , R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{15} , R^{16} , A^2 and G^{11} are as described above, with an acid;

10 10. a production method of a phosphine compound of formula (24B) :



wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{15} , R^{16} , A^2 and G^{11} are as described above,

which comprises reacting
a phosphine compound of formula (24C):



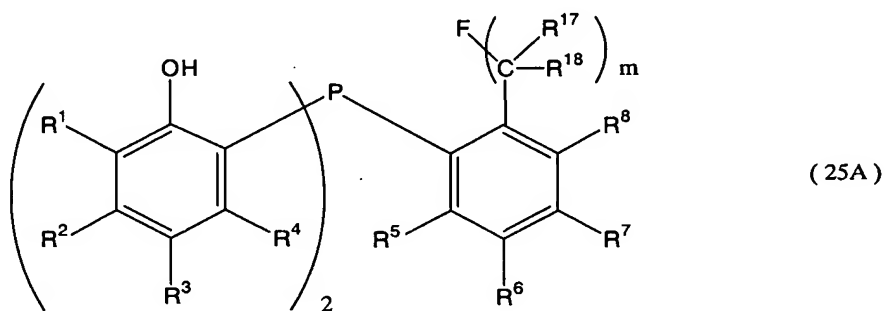
wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{15} , A^2 and G^{11} are
5 as described above,

with a metal hydride compound or a metal aryl compound
of formula (24D);



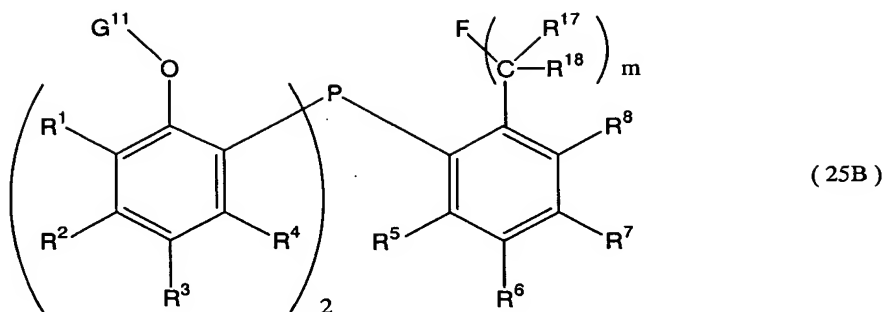
wherein R^{16} and Y represent an alkali metal, or
10 $J-X^3$, wherein J represents an alkali earth metal and X^3
represents a halogen atom;

11. a production method of a phosphine compound of formula
(25A):



15 wherein , R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{17} , R^{18} and m are
as described above,

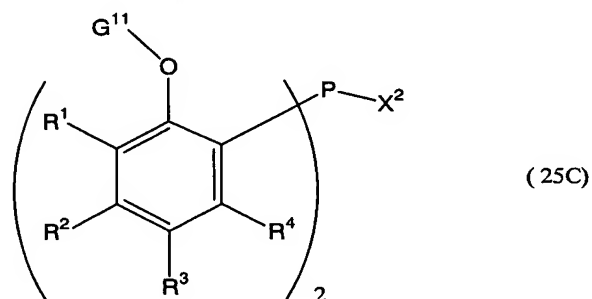
which comprises reacting
a phosphine compound of formula (25B):



wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{17} , R^{18} , G^{11} and m are as described above,

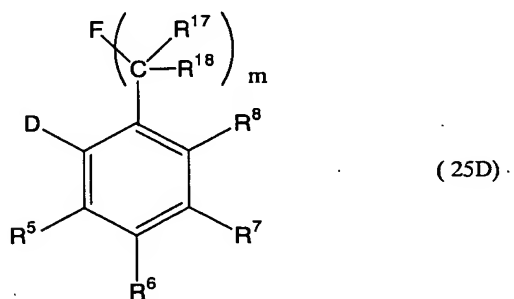
with an acid;

- 5 12. a production method of a phosphine compound of formula (25B) above, which comprises reacting a phosphine halide compound of formula (25C):



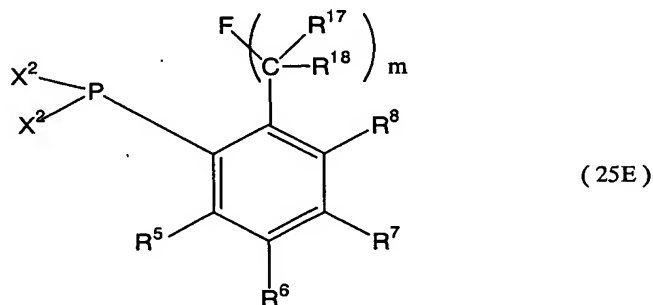
- 10 wherein R^1 , R^2 , R^3 , R^4 and G^{11} are as above, and X^2 represented a halogen atom,

with a metal aryl compound of formula (25D);

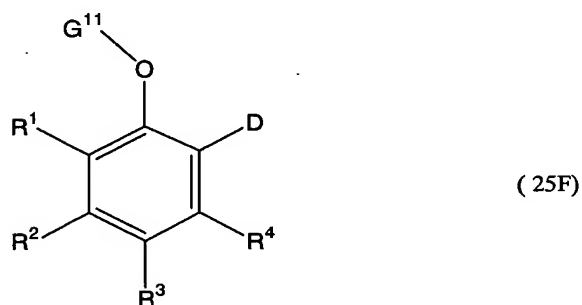


wherein R^5 , R^6 , R^7 , R^8 , R^{15} , R^{16} , D and m are as described above;

13. a production method of a phosphine compound of formula (25B) above, which comprises reacting a halo-phosphine compound of formula (25E):

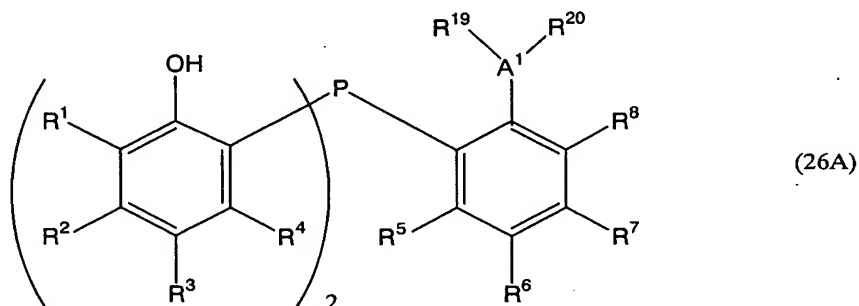


5 wherein R^5 , R^6 , R^7 , R^8 , R^{95} , R^{96} and m are as described above, and X^2 represents a halogen atom, with a metal aryl compound of formula (25F);



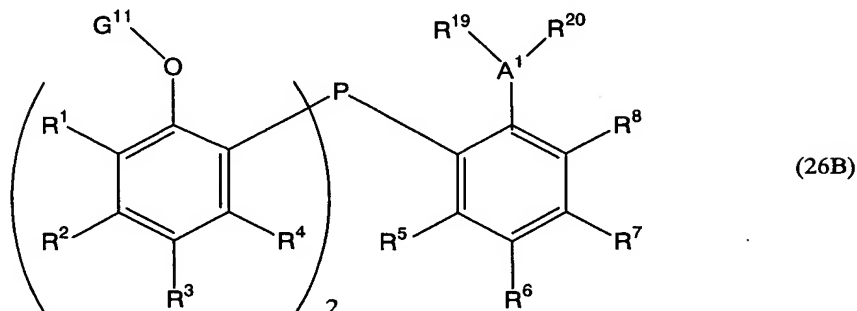
wherein R^1 , R^2 , R^3 , R^4 , G^{11} and D are as described above;

10 14. a production method of a phosphine compound of formula (26A):



wherein A^1 , R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{19} , R^{20} and A^1 are as described above,

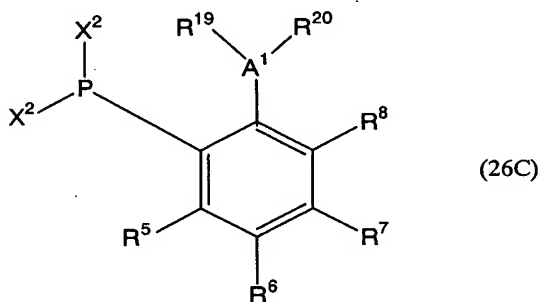
which comprises reacting
a phosphine compound of formula (26B):



wherein A^1 , R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{19} , R^{20} and G^{11}
5 are as described above,

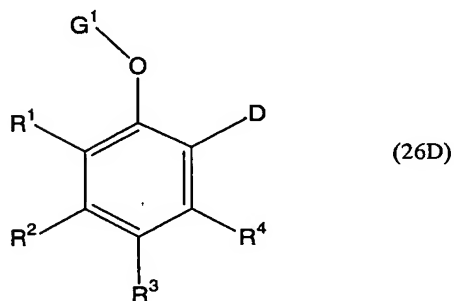
with an acid;

15. a production method of a phosphine compound of formula
(26B) above, which comprises reacting
a halo-phosphine compound of formula (26C):



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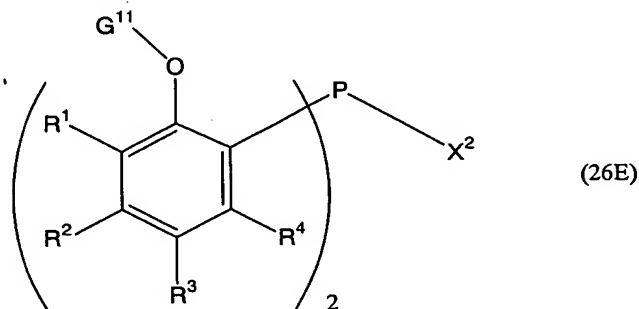
wherein R^5 , R^6 , R^7 , R^8 , R^{19} , R^{20} and A^1 are as described above,
and X^2 represents a halogen atom,
with a metal aryl compound of formula (26D):



wherein R^1 , R^2 , R^3 , R^4 , D and G^1 (preferably G^{11}) are as described above;

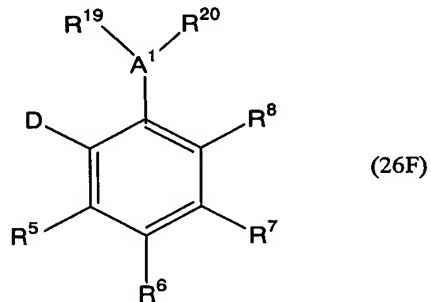
16. a production method of a phosphine compound of formula (26B) above, which comprises reacting

5 an aryl halogenated phosphorous compound of formula (26E):



wherein R^1 , R^2 , R^3 , R^4 and G^{11} are as described above, and X^2 represents a halogen atom,

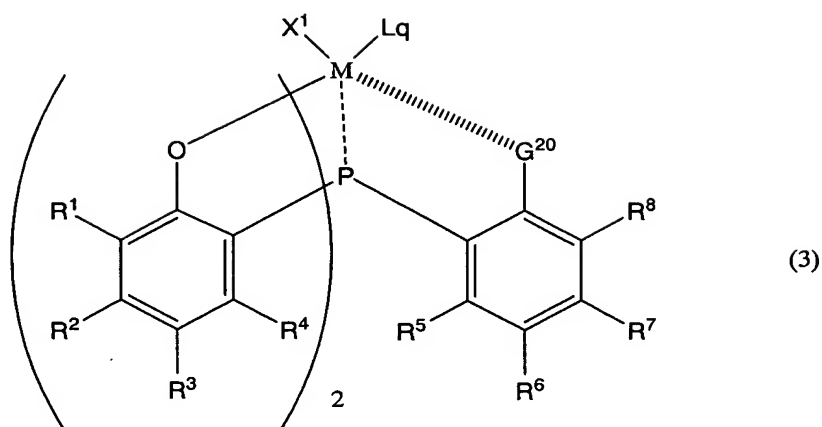
with a metal aryl compound of formula (26F):



10

wherein A^1 , R^5 , R^6 , R^7 , R^8 , R^{19} , R^{20} and D are as described above;

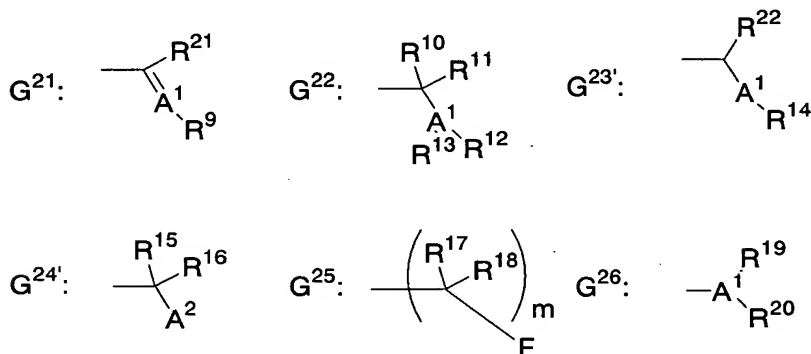
17. a production method of a transition metal complex of formula (3):



wherein M, R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, X¹ and L are as described below,

q represents an integer of 0 or 1;

5 G²⁰ represents any one of G²¹ to G²⁶ below;



wherein A¹ represents an element of Group 15 of the periodic table, A¹ in G^{23'} represents an anion of an element of Group 15 of the periodic table, A² represents an anion of an element of Group 16 of the periodic table, and A¹ in G²¹ represents a nitrogen atom;

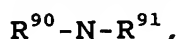
R⁹ and R¹⁴ each represents,

a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

15 a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

a substituted or unsubstituted aryl group having 6 to 20 carbon atoms, or

a group of formula:



5 wherein R^{90} and R^{91} are the same or different, and represent a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms,

10 a substituted or unsubstituted aryl group having 6 to 20 carbon atoms, or

a cyclic structure being linked together;

R^{12} , R^{13} , R^{19} and R^{20} each independently represents,

a substituted or unsubstituted alkyl group having 1 to 15 10 carbon atom(s),

a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms, or

a substituted or unsubstituted aryl group having 6 to 20 carbon atoms; or

20 R^{12} and R^{13} , and R^{19} and R^{20} each independently represent a cyclic group being linked together;

R^{10} , R^{11} , R^{15} , R^{16} , R^{21} and R^{22} each independently represents,

a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

25 a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms, or

a substituted or unsubstituted aryl group having 6 to 20 carbon atoms;

R^{17} and R^{18} are the same or different, and each represents

a hydrogen atom,

a halogen atom,

a substituted or unsubstituted alkyl group having 1 to 10 carbon atom(s),

5 a substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms, or

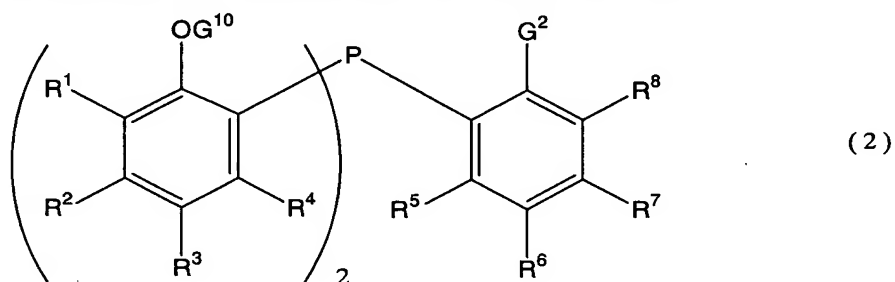
a substituted or unsubstituted aryl group having 6 to 20 carbon atoms;

m represents an integer of 0 or 1; and

10 the line linking M and G²⁰ means that M is linked or coordinated to an element of Group 15 or 16 of the periodic table or to a fluorine atom constituting G²⁰,

which comprises reacting

a phosphine compound of formula (2):



15 wherein R¹, R², R³, R⁴, R⁵, R⁶, R⁷ and R⁸ are as described above, and G¹⁰ represents

a hydrogen atom, or a protective group of hydroxyl group selected from an alkyl group having a secondary or tertiary carbon atom linked to an oxygen atom of phenol or a C1 to C2 alkyl group substituted with a substituted or unsubstituted alkoxy group,

with a transition metal compound of formula (4):



wherein

25 M represents an element of Group 4 of the periodic table;

X^1 represents,

a hydrogen atom,

a halogen atom,

a substituted or unsubstituted alkyl group having 1 to
5 10 carbon atom(s),

a substituted or unsubstituted aralkyl group having 7 to
20 carbon atoms,

a substituted or unsubstituted aryl group having 6 to 20
carbon atoms,

10 a substituted or unsubstituted alkoxy group having 1 to
10 carbon atom(s),

a substituted or unsubstituted aralkyloxy group having
7 to 20 carbon atoms,

a substituted or unsubstituted aryloxy group having 6 to
15 20 carbon atoms, or

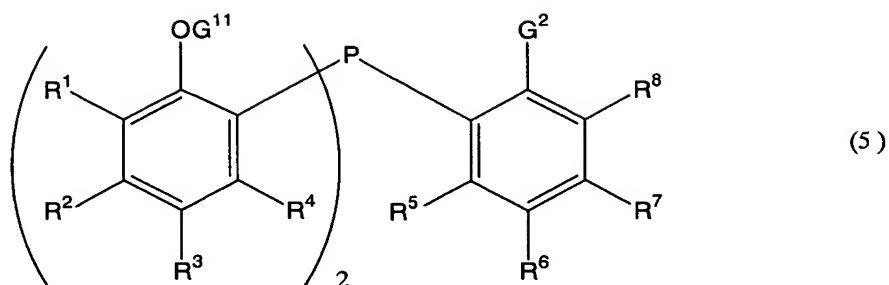
an amino group disubstituted with a hydrocarbon having
2 to 20 carbon atoms;

L represents a balancing counter ion or neutral ligand,
being an atom or a group similar to X^1 , and is bonding or
20 coordinating to metal M, L^1 represents a neutral ligand, and
p represents an integer of 0 to 2;

18. a production method of the transition metal complex
of formula (3) above,

which comprises reacting

25 a phosphine compound of formula (5):



wherein R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , G^{11} and G^2 are as described above,

with the transition metal compound of formula (4) above;

5 19. the transition metal complex of formula (3);

20. an olefin polymerization catalyst prepared by combining the transition metal compound of formula (3) and the following compound A, and optionally compound B;

10 compound A: any one of A1 to A3 below, or a mixture of at least two of them;

A1: an organic aluminum compound of formula $(E1)_aAl(Z)_{(3-a)}$;

A2: a cyclic aluminosiloxane having a structure represented by $[-Al(E2)-O-]_b$; and

15 A3: a linear aluminosiloxane having a structure represented by $(E3)[-Al(E3)-O-]_cAl(E3)_2$;

20 wherein , $E1$ to $E3$ are the same or different and are hydrocarbons having 1 to 8 carbon atom(s); Z is the same or different and represents a hydrogen atom or a halogen atom; a represents 1, 2 or 3; b represents an integer of 2 or more; and c represents an integer of 1 or more; and

compound B: any one of B1 to B3 below, or a mixture of at least two of them;

B1: a boron compound of formula $BQ^1Q^2Q^3$;

B2: a boron compound of formula $Z^+(BQ^1Q^2Q^3Q^4)$; and

25 B3: a boron compound of formula $(L-H)^+(BQ^1Q^2Q^3Q^4)$;

wherein B is a boron atom in a trivalent atomic valance state, Q^1 to Q^4 are the same or different, and represent a halogen atom, a hydrocarbon having 1 to 20 carbon atom(s), a halogenated hydrocarbon having 1 to 20 carbon atom(s), an alkoxy group having
5 1 to 20 carbon atom(s), a silyl group substituted with a hydrocarbon having 1 to 20 carbon atom(s), or an amino group disubstituted with hydrocarbons having 1 to 20 carbon atom(s); and

21. a production method of an olefin polymer for
10 polymerizing an olefin using the olefin polymerization catalyst above.

Best Mode for Carrying Out the Invention

Substituents of the phosphine compound of formula (1) will be described below.

15 Specific examples of the halogen atom represented by R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^{17} or R^{18} include a fluorine atom, a chlorine atom, a bromine atom and an iodine atom, and the chlorine atom is preferable as the halogen atoms of R^1 , R^2 , R^3 , R^4 , R^6 , R^7 and R^8 , and the fluorine atom is preferable as the halogen
20 atoms of R^{17} and R^{18} .

Specific examples of the unsubstituted C1 to C10 alkyl groups represented by R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{90} or R^{91} include methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl,
25 tert-butyl, n-pentyl, neopentyl, amyl, n-hexyl, n-octyl and n-decyl groups. Examples of the substituted C1 to C10 alkyl groups include the C1 to C10 alkyl groups substituted with a halogen atom, an alkoxy group, an aryloxy group, a hydrocarbon-substituted amino group or a silyl group substituted

with a hydrocarbon, and specific examples thereof include a fluoromethyl group, a difluoroethyl group, a trifluoromethyl group, a fluoroethyl group, a difluoroethyl group, a trifluoroethyl group, a tetrafluoroethyl group, a pentafluoroethyl group, a perfluoropropyl group, a perfluorobutyl group, a perfluoropentyl group, a perfluorohexyl group, a perfluorooctyl group, a perfluorodecyl group, a trichloromethyl group, a methoxymethyl group, a phenoxymethyl group, a diaminomethyl group and a trimethylsilylmethyl group.

Among the alkyl groups that may be substituted having 1 to 10 carbon atom(s), the methyl, ethyl, isopropyl, tert-butyl and amyl groups are preferable, and the methyl and tert-butyl groups are more preferable.

Examples of unsubstituted C7 to C20 aralkyl groups represented by R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{90} and R^{91} include benzyl, naphthylmethyl, anthracenylmethyl and diphenylmethyl groups; examples of the substituted C7 to C20 aralkyl groups include those substituted with a halogen atom, an alkyl group, an alkoxy group, an aryloxy group or a hydrocarbon-substituted amino group, or with a silyl group substituted with a hydrocarbon; and specific examples of the substituted or unsubstituted aralkyl group having 7 to 20 carbon atoms include (2-methylphenyl)methyl group, (3-methylphenyl)methyl group, (4-methylphenyl)methyl group, (2,3-dimethylphenyl)methyl group, (2,4-dimethylphenyl)methyl group, (2,5-dimethylphenyl)methyl group, (2,6-dimethylphenyl)methyl group, (3,4-dimethylphenyl)methyl group, (2,3,4-trimethylphenyl)methyl group,

(2,3,5-trimethylphenyl)methyl group,
 (2,3,6-trimethylphenyl)methyl group,
 (3,4,5-trimethylphenyl)methyl group,
 (2,4,6-trimethylphenyl)methyl group,
 5 (2,3,4,5-tetramethylphenyl)methyl group,
 (2,3,4,6-tetramethylphenyl)methyl group,
 (2,3,5,6-tetramethylphenyl)methyl group,
 (pentamethylphenyl)methyl group, (ethylphenyl)methyl group,
 (n-propylphenyl)methyl group, (isopropylphenyl)methyl group,
 10 (n-butylphenyl)methyl group, (sec-butylphenyl)methyl group,
 (tert-butylphenyl)methyl group, (n-pentylphenyl)methyl group,
 (neopentylphenyl)methyl group, (n-hexylphenyl)methyl group,
 (n-octylphenyl)methyl group, (n-decylphenyl)methyl group,
 (n-dodecylphenyl)methyl group, (fluorophenyl)methyl group,
 15 (difluorophenyl)methyl group, (pentafluorophenyl)methyl group,
 (chlorophenyl)methyl group, (phenoxyphenyl)methyl group,
 (dimethylaminophenyl)methyl group and
 (trimethylsilylphenyl)methyl group. The benzyl group is
 preferable as the substituted or unsubstituted aralkyl group
 20 having 7 to 20 carbon atoms.

Examples of unsubstituted C₆ to C₂₀ aryl groups represented
 by R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹, R¹², R¹³, R¹⁴, R¹⁵,
 R¹⁶, R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R⁹⁰ or R⁹¹ include phenyl, naphthyl
 and anthracenyl groups, and examples of substituted C₆ to C₂₀
 25 aryl groups include those substituted with a halogen atom, an
 alkyl group, an alkoxy group, an aryloxy group and a
 hydrocarbon-substituted amino group, or with a silyl group
 substituted with a hydrocarbon. Specific examples thereof
 include 2-tolyl, 3-tolyl, 4-tolyl, 2,3-xylyl, 2,4-xylyl,

2,5-xylyl, 2,6-xylyl, 3,4-xylyl, 3,5-xylyl,
 2,3,4-trimethylphenyl, 2,3,5-trimethylphenyl,
 2,3,6-trimethylphenyl, 2,4,6-trimethylphenyl,
 3,4,5-trimethylphenyl, 2,3,4,5-tetramethylphenyl,
 5 2,3,4,6-tetramethylphenyl, 2,3,5,6-tetramethylphenyl,
 pentamethylphenyl, ethylphenyl, n-propylphenyl,
 isopropylphenyl, n-butylphenyl, sec-butylphenyl,
 tert-butylphenyl, n-pentylphenyl, neopentylphenyl,
 n-hexylphenyl, n-octylphenyl, n-decylphenyl, n-dodecylphenyl,
 10 n-tetradecylphenyl, 2-fluorophenyl, 3-fluorophenyl,
 4-fluorophenyl, 3,5-difluorophenyl, pentafluorophenyl,
 4-chlorophenyl, 2-methoxyphenyl, 3-methoxyphenyl,
 4-methoxyphenyl, 4-phenoxyphenyl, 4-dimethylaminophenyl and
 4-trimethylsilylphenyl groups. Preferred substituted or
 15 unsubstituted C₆ to C₂₀ aryl group is the phenyl group.

Examples of cyclic structures formed by linking R¹² and
 R¹³ together include C₄ to C₆ cyclic alkylene groups such as
 tetramethylene, pentamethylene and hexamethylene groups.
 Examples of cyclic structures formed by linking R¹⁹ and R²⁰ and
 20 R⁹⁰ and R⁹¹ together include the same groups. Examples of cyclic
 structures formed by linking R⁹⁰ and R⁹¹ together include 1-pyrrolyl
 group.

Examples of the hydrocarbon of the silyl group substituted
 with unsubstituted C₁ to C₂₀ hydrocarbons represented by R¹,
 25 R², R³, R⁴, R⁵, R⁶, R⁷ or R⁸ include alkyl groups having 1 to 10
 carbon atom(s) such as methyl, ethyl, n-propyl, isopropyl,
 n-butyl, sec-butyl, tert-butyl, n-pentyl, neopentyl, amyl,
 n-hexyl, cyclohexyl, n-octyl and n-dodecyl groups; and C₆ to
 C₂₀ aryl groups such as phenyl, tolyl, xylyl, naphthyl and

anthracenyl groups. Examples of the silyl group substituted with these C1 to C20 hydrocarbons include monosubstituted silyl groups such as methylsilyl, ethylsilyl, phenylsilyl group; disubstituted silyl groups such as dimethylsilyl, diethylsilyl or diphneysilyl group; and tri-substituted silyl groups such as trimethylsilyl, triethylsilyl, tri-n-propylsilyl, tri-isopropylsilyl, tri-n-butylsilyl, tri-sec-butylsilyl, tri-tert-butylsilyl, tri-isobutylsilyl, tert-butyl dimethylsilyl, tri-n-pentylsilyl, tri-n-pentylsilyl, tri-n-hexylsilyl, tricyclohexylsilyl or triphenylsilyl group. Examples of the preferable silyl group substituted with substituted or unsubstituted C1 to C20 hydrocarbons include trimethylsilyl, tert-butyl dimethyl and triphenylsilyl groups. Examples of the silyl group substituted with substituted C1 to C20 hydrocarbons include all the silyl groups substituted with such a hydrocarbon that is substituted with a halogen atom, for example, a fluorine atom.

Specific examples of unsubstituted C1 to C10 alkoxy groups represented by R^1 , R^2 , R^3 , R^4 , R^6 , R^7 or R^8 include methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, sec-butoxy, tert-butoxy, n-pentyloxy, neopentyloxy, n-hexyloxy, n-octyloxy, n-nonyloxy and n-decyloxy groups. Examples of substituted C1 to C10 alkoxy groups include C1 to C10 alkoxy groups substituted with a halogen atom, an alkoxy group, an aryloxy group, a hydrocarbon-substituted amino group, or with a silyl group substituted with a hydrocarbon.

Specific examples of the substituted alkoxy group include fluoromethoxy, difluoromethoxy, trifluoromethoxy, fluoroethoxy, difluoroethoxy, trifluoroethoxy,

tetrafluoroethoxy, pentafluoroethoxy, perfluoropropoxy,
perfluorobutyloxy, perfluoropentyloxy, perfluorohexyloxy,
perfluorooctylcoxy, perfluorodecyloxy, trichloromethyloxy,
methoxymethoxy, phenoxymethoxy, dimethylaminomethoxy and
5 trimethylsilylmethoxy groups. Preferred are substituted or
unsubstituted C1 to C10 alkoxy groups such as methoxy, ethoxy,
tert-butoxy group or the like.

Examples of unsubstituted C7 to C20 aralkyloxy groups
represented by R^1 , R^2 , R^3 , R^4 , R^7 or R^8 include benzyloxy,
10 naphthylmethoxy, anthracenylmethoxy and diphenylmethoxy
groups.

Examples of the substituted C7 to C20 aralkyloxy group
include those substituted with a halogen atom, an alkyl group,
an alkoxy group, an aryloxy group, a hydrocarbon-substituted
15 aminogroup, or with a silyl group substituted with a hydrocarbon.
Specific examples thereof include (2-methylphenyl)methoxy
group, (3-methylphenyl)methoxy group, (4-methylphenyl)methoxy
group, (2,3-dimethylphenyl)methoxy group,
(2,4-dimethylphenyl)methoxy group,
20 (2,5-dimethylphenyl)methoxy group,
(2,6-dimethylphenyl)methoxy group,
(3,4-dimethylphenyl)methoxy group,
(2,3,4-trimethylphenyl)methoxy group,
(2,3,5-trimethylphenyl)methoxy group,
25 (2,3,6-trimethylphenyl)methoxy group,
(3,4,5-trimethylphenyl)methoxy group,
(2,4,6-trimethylphenyl)methoxy group,
(2,3,4,5-tetramethylphenyl)methoxy group,
(2,3,4,6-tetramethylphenyl)methoxy group,

(2,3,5,6-tetramethylphenyl)methoxy group,
(pentamethylphenyl)methoxy group, (ethylphenyl)methoxy group,
(n-propylphenyl)methoxy group, (isopropylphenyl)methoxy group,
(n-butylphenyl)methoxy group, (sec-butylphenyl)methoxy group,
5 (tert-butylphenyl)methoxy group, (n-pentylphenyl)methoxy
group, (neopentylphenyl)methoxy group, (n-hexylphenyl)methoxy
group, (n-octylphenyl)methoxy group, (n-decylphenyl)methoxy
group, (n-dodecylphenyl)methoxy group, (fluorophenyl)methyl
group, (difluorophenyl)methyl group,
10 (pentafluorophenyl)methyl group, (chlorophenyl)methyl group,
(methoxyphenyl)methyl group, (phenoxyphenyl)methyl group,
(dimethylaminophenyl)methyl group and
(trimethylsilylphenyl)methyl group. Preferable examples of
the substituted or unsubstituted C7 to C20 aralkyloxy groups
15 include benzyloxy group and the like.

Examples of the unsubstituted or substituted C6 to C20
aryloxy group represented by R^1 , R^2 , R^3 , R^4 , R^7 or R^8 include phenoxy,
naphthoxy and anthracenoxy groups.

Examples of the substituted C6 to C20 aryloxy group include
20 those substituted with a halogen atom, an alkyl group, an alkoxy
group, an aryloxy group, a hydrocarbon-substituted amino group,
or with a silyl group substituted with a hydrocarbon.

Specific examples thereof include 2-methylphenoxy group,
3-methylphenoxy group, 4-methylphenoxy group,
25 2,3-dimethylphenoxy group, 2,4-dimethylphenoxy group,
2,5-dimethylphenoxy group, 2,6-dimethylphenoxy group,
3,4-dimethylphenoxy group, 3,5-dimethylphenoxy group,
2,3,4-trimethylphenoxy group, 2,3,5-trimethylphenoxy group,
2,3,6-trimethylphenoxy group, 2,4,5-trimethylphenoxy group,

2,4,6-trimethylphenoxy group, 3,4,5-trimethylphenoxy group,
 2,3,4,5-tetramethylphenoxy group, 2,3,4,6-tetramethylphenoxy
 group, 2,3,5,6-tetramethylphenoxy group, pentamethylphenoxy
 group, ethylphenoxy group, n-propylphenoxy group,
 5 isopropylphenoxy group, n-butylphenoxy group, sec-butylphenoxy
 group, tert-butylphenoxy group, n-hexylphenoxy group,
 n-octylphenoxy group, n-decylphenoxy group,
 n-tetradecylphenoxy group, 2-fluorophenoxy group,
 3-fluorophenoxy group, 4-fluorophenoxy group,
 10 3,5-difluorophenoxy group, pentafluorophenoxy group,
 4-chlorophenoxy group, 2-methoxyphenoxy group, 3-methoxyphenoxy
 group, 4-methoxyphenoxy group, 4-phenoxyphenoxy group,
 4-dimethylaminophenoxy group and 4-trimethylsilylphenoxy group.
 Preferable examples of the substituted or unsubstituted C7 to
 15 C20 aryloxy groups include the phenoxy group and the like.

The amino group disubstituted with the unsubstituted C1
 to C20 hydrocarbon, represented by R^1 , R^2 , R^3 , R^4 , R^6 , R^7 or R^8 ,
 is an amino group substituted with two hydrocarbons, wherein
 examples of the hydrocarbon include C1 to C10 alkyl groups such
 20 as methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl,
 tert-butyl, n-pentyl, neopentyl, amyl, n-hexyl, cyclohexyl,
 n-octyl or n-decyl; and C6 to C20 aryl groups such as phenyl,
 tolyl, xylyl, naphthyl or anthracenyl group. Examples of the
 amino group substituted with these C1 to C20 hydrocarbons include
 25 dimethylamino group, diethylamino group, di-n-propylamino
 group, di-isopropylamino group, di-n-butylamino group,
 di-sec-butylamino group, di-tert-butylamino group,
 di-isobutylamino group, tert-butylisopropylamino group,
 di-n-hexylamino group, di-n-octylamino group, di-n-decylamino

group and diphenylamino group, and preferable examples include dimethylamino group and diethylamino group.

Examples of the protective group represented by G^1 include protective groups of hydroxyl group selected from alkyl groups
5 having a secondary or tertiary carbon atom linked to an oxygen atom of phenol, or C1 to C2 alkyl groups substituted with a substituted or unsubstituted alkoxy group, or a substituted or unsubstituted C7 to C20 aralkyl groups.

Examples of the alkyl group having secondary or tertiary
10 carbon atom linked to the oxygen atom of phenol include isopropyl, tert-butyl and sec-butyl groups.

Examples of the protective group of the hydroxyl group selected from C1 to C2 alkyl groups substituted with the substituted or unsubstituted alkoxy group include methoxymethyl,
15 ethoxyethyl, methoxyethoxymethyl trimethylsilylethoxymethyl or 1-ethoxyethyl group.

Examples of the substituted or unsubstituted C7 to C20 aralkyl groups include those as described above. Examples of the preferable protective group include the protective group
20 of the hydroxyl group, as shown by G^{11} , selected from alkyl groups having the secondary or tertiary carbon atom linked to the oxygen atom of phenol, and C1 to C2 alkyl groups substituted with a substituted or unsubstituted alkoxy group, and examples of the more preferable protective group include methoxymethyl,
25 ethoxyethyl, methoxyethoxymethyl, trimethylsilylethoxymethyl or 1-ethoxyethyl group that is a protective group of the hydroxyl group selected from C1 to C2 alkyl groups substituted with substituted or unsubstituted alkoxy group.

In G^{21} , G^{22} , G^{23} , $G^{23'}$ and G^{26} of G^2 , the 15th group of the

periodic table represented by A^1 is preferably a nitrogen atom, and the 16th group of the periodic table represented by A^2 in G^{24} and $G^{24'}$ is preferably an oxygen atom.

5 Examples of the compound in which G^1 represents a hydrogen atom in formula (1) include the compounds of formulae 21A, 22A, 23A, 24A, 25A and 26A. Examples of the compound representing G^{11} in which G^1 represents a group other than hydrogen include the compounds of formulae 21B, 22B, 23B, 24B and 25B.

10 The compound of formula (21A) can be produced by reacting the compound of formula (21B) with the organic compound of formula 21F.

The compound of formula (21A) can be obtained by reacting the compound of formula (21B) with an acid, wherein the compound of formula (21B) is a protective group of the hydroxyl group
15 selected from alkyl groups having a secondary or tertiary carbon atom linked to the oxygen atom of phenol, and C1 to C2 alkyl groups substituted with a substituted or unsubstituted alkoxy group. Examples of the production method of the compounds of formulae (21B) and (21A), in which G^{11} is the protective group
20 of the hydroxyl group selected from alkyl groups having a secondary or tertiary carbon atom linked to the oxygen atom of phenol, and C1 to C2 alkyl groups substituted with a substituted or unsubstituted alkoxy group, will be described below. The compound of formula (22A), (23A), (24A) or (25A) may be also
25 produced in a similar manner above by a reaction with the corresponding compound of formula (22B), (23B), (24B) or (25B), respectively.

An example of the acid in this reaction is a Bronsted acid (for example, inorganic acids), and detailed examples thereof

include halogenated hydrogen such as hydrogen chloride, hydrogen bromide and hydrogen iodide, and sulfuric acid, preferably hydrogen chloride. The compound of formula (25A) may be obtained from the compound of formula (22A) by deprotection with an acid
5 as a salt of Bronsted acid (for example, salts of inorganic acids or halogenated hydrogen) such as phosphine, amine and imine salts.

Hydrogen chloride gas may be used as the hydrogen chloride used in the reaction above, or hydrogen chloride may be generated
10 from an acid chloride and an alcohol in the reaction system.

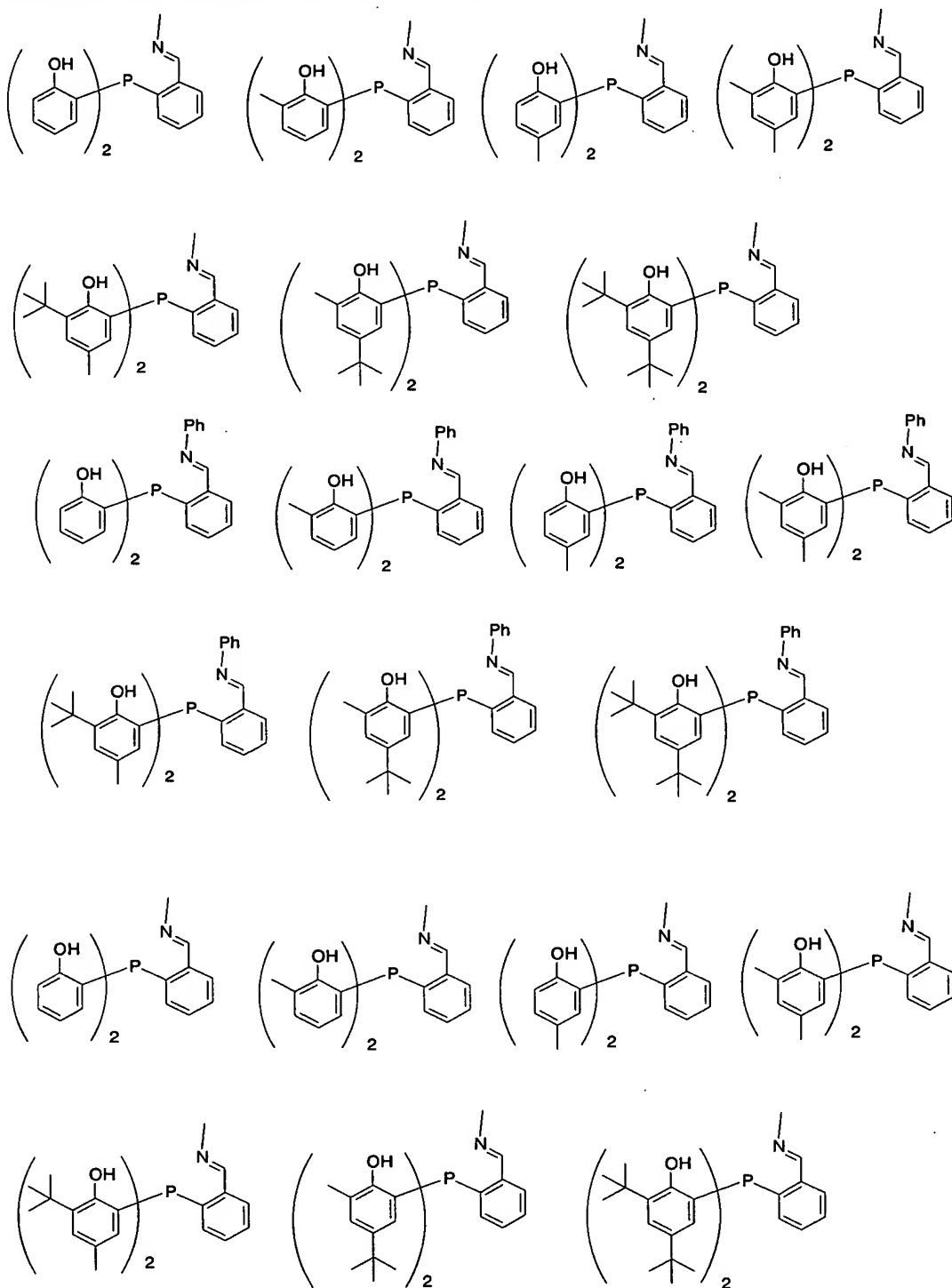
The reaction is usually carried out in a solvent inert to the reaction. Examples of the solvent available are aprotic solvents including aromatic hydrocarbon solvents such as benzene or toluene; aliphatic hydrocarbon solvents such as hexane or
15 heptane; ether solvents such as diethyl ether, tetrahydrofuran or 1,4-dioxane; polar solvents such as acetonitrile, propionitrile, acetone, diethyl ketone, methyl isobutyl ketone, cyclohexanone or ethyl acetate; and halogenated solvents such as dichloromethane, dichloroethane, chlorobenzene or
20 dichlorobenzene. These solvents may be used alone or as a mixture of at least two of them, and the amount thereof is usually in the range of 1 to 200 parts by weight, preferably 3 to 50 parts by weight per part by weight of the phosphine compound of formula (21B).

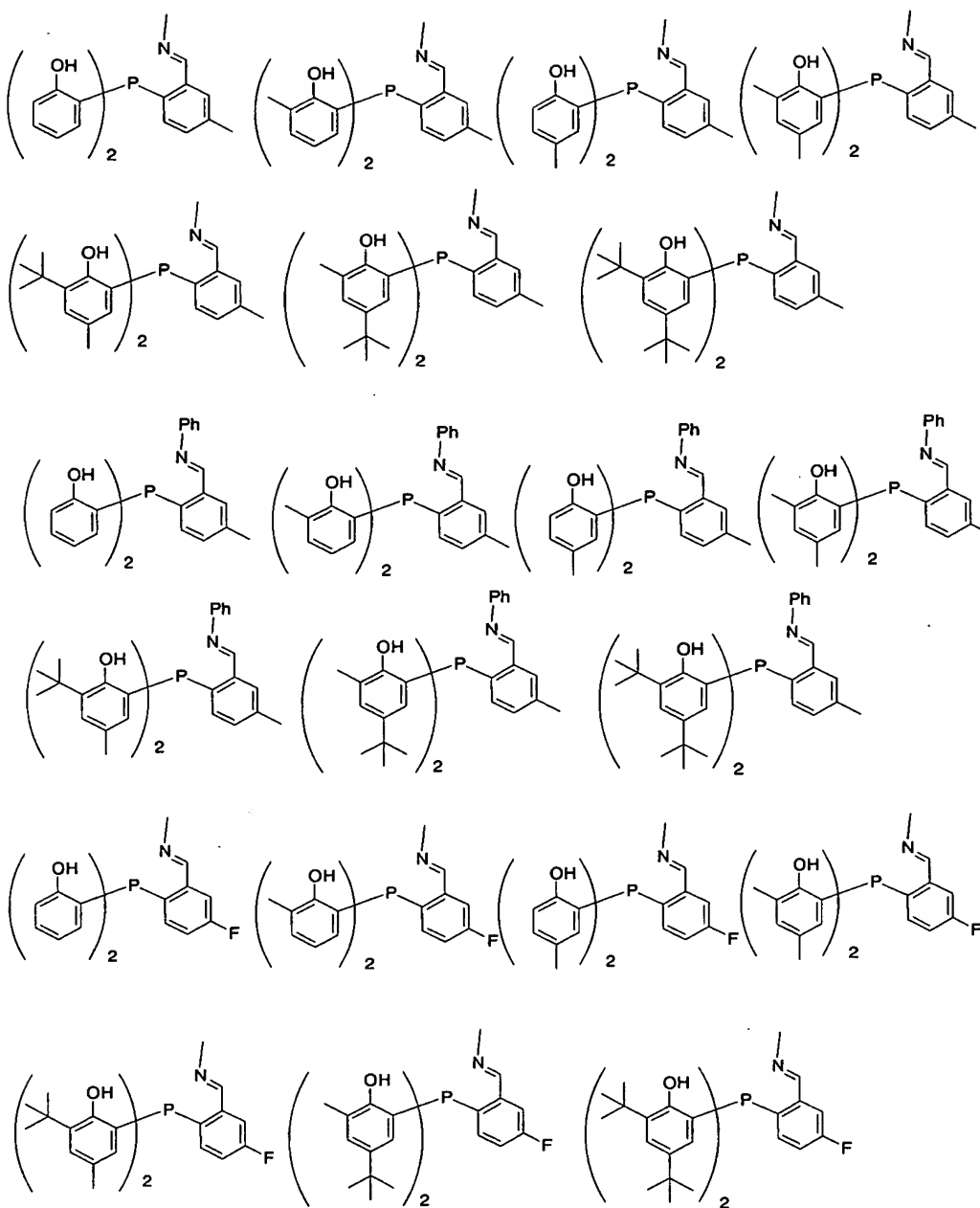
25 The reaction temperature is usually in the range of -100°C or more to the boiling point or less of the solvent, preferably about -80 to 100°C .

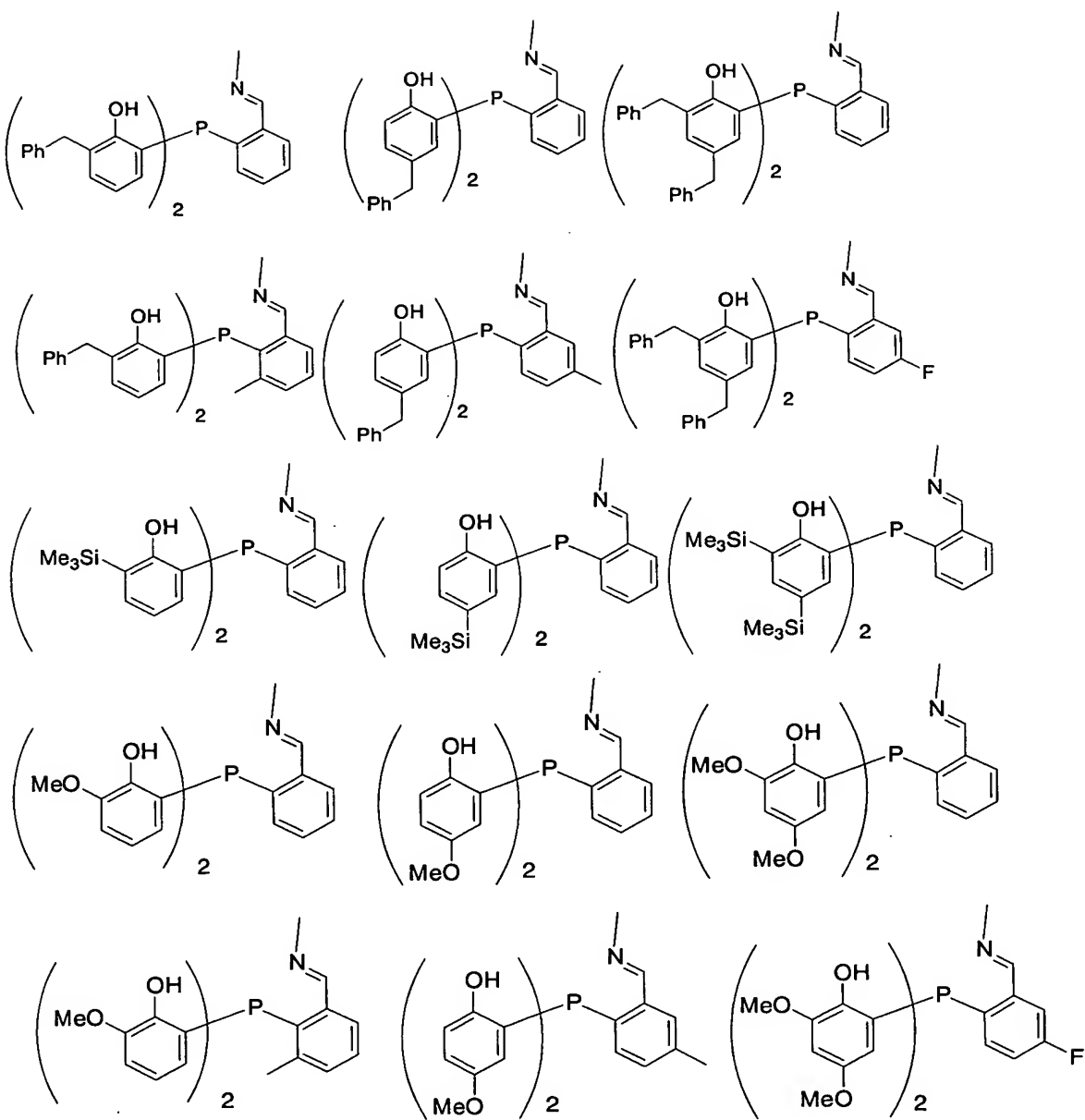
The phosphine compound of formula (22A) can be obtained by a conventional method for obtaining a product from a reaction

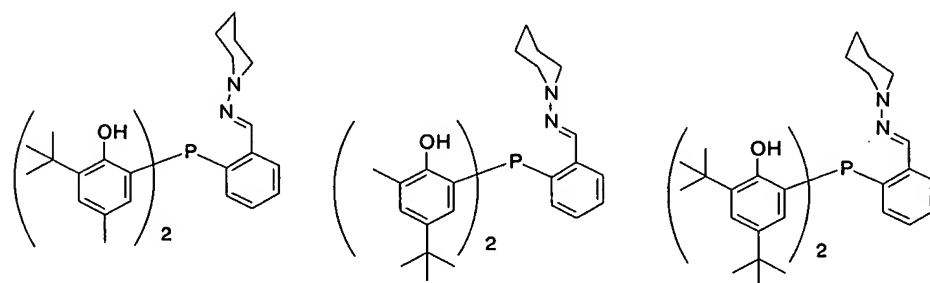
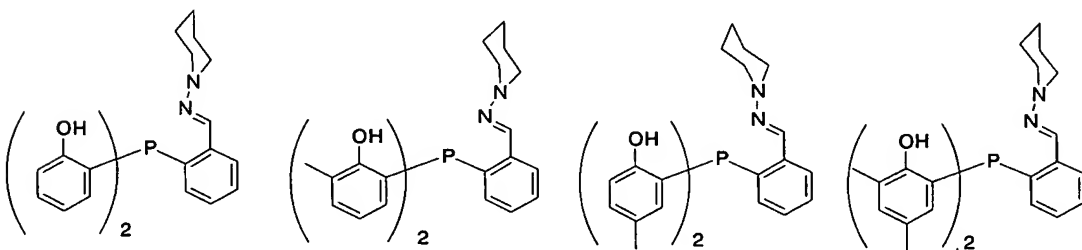
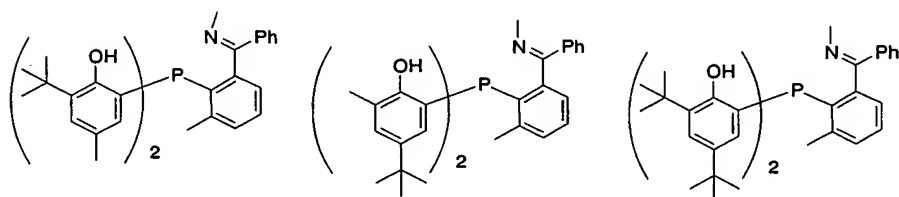
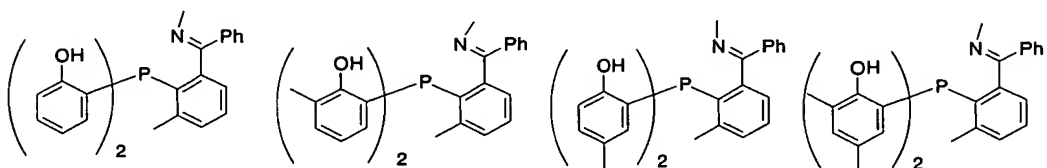
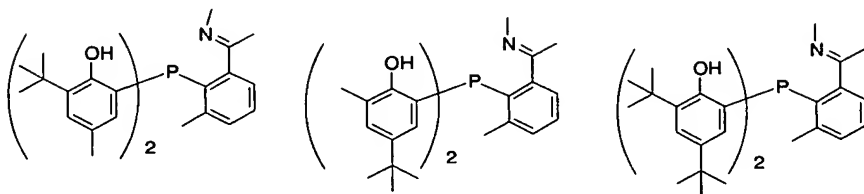
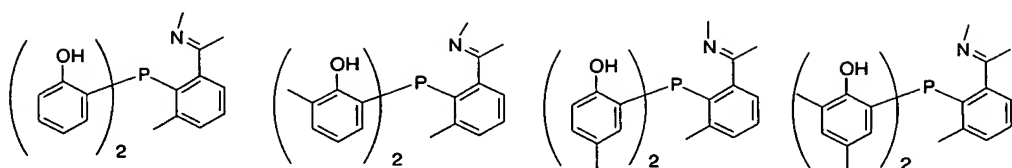
mixture, for example by removing the solvent by evaporation.

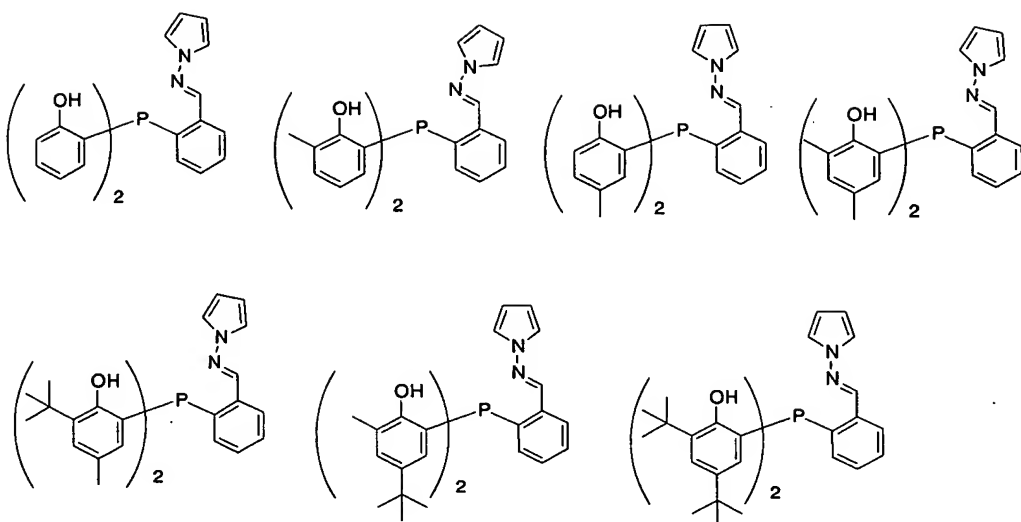
The following compounds are the specific examples of the phosphine compounds of formula (21A):



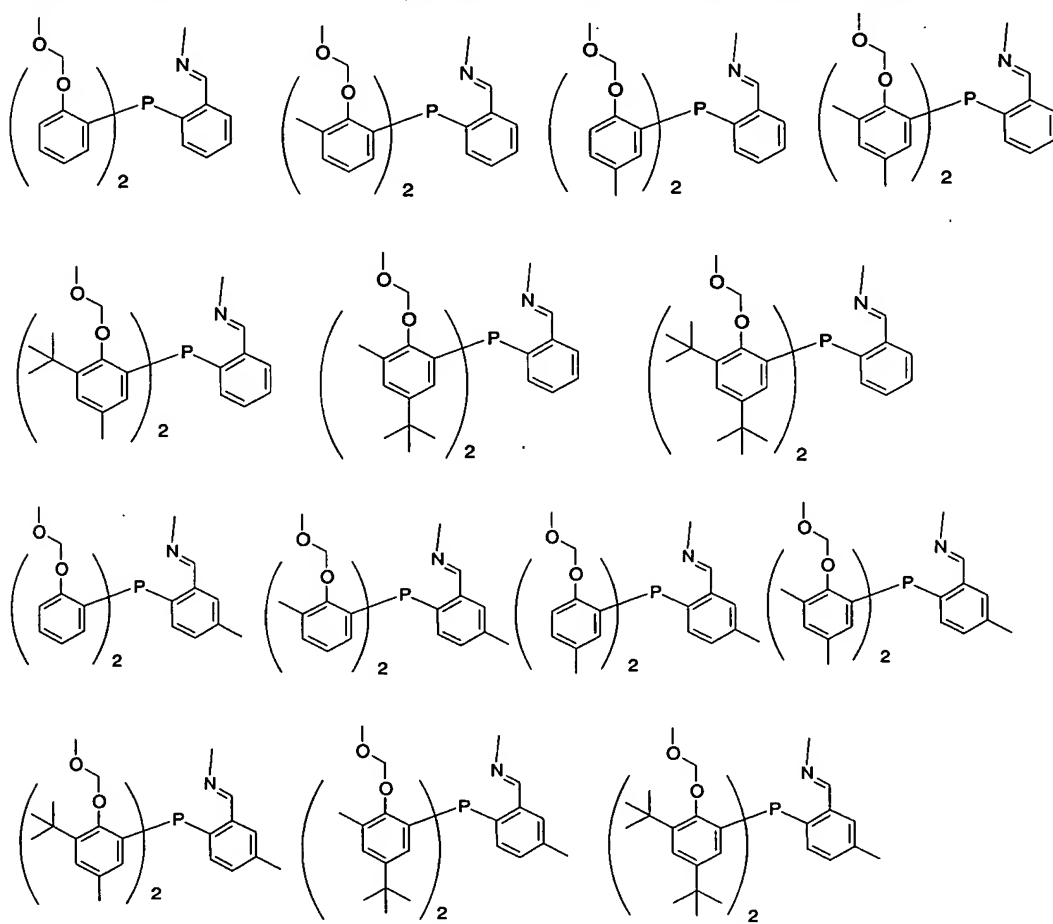


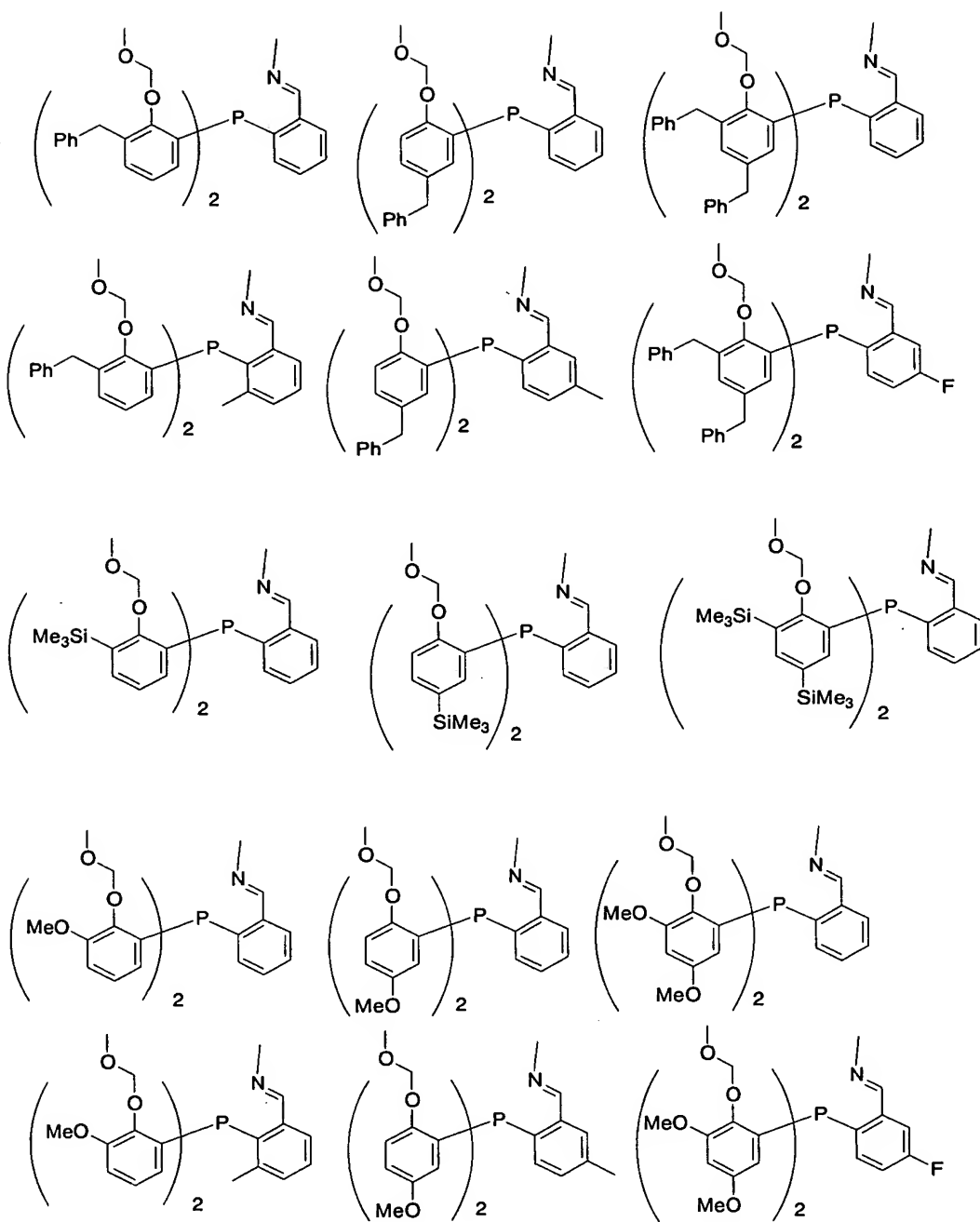




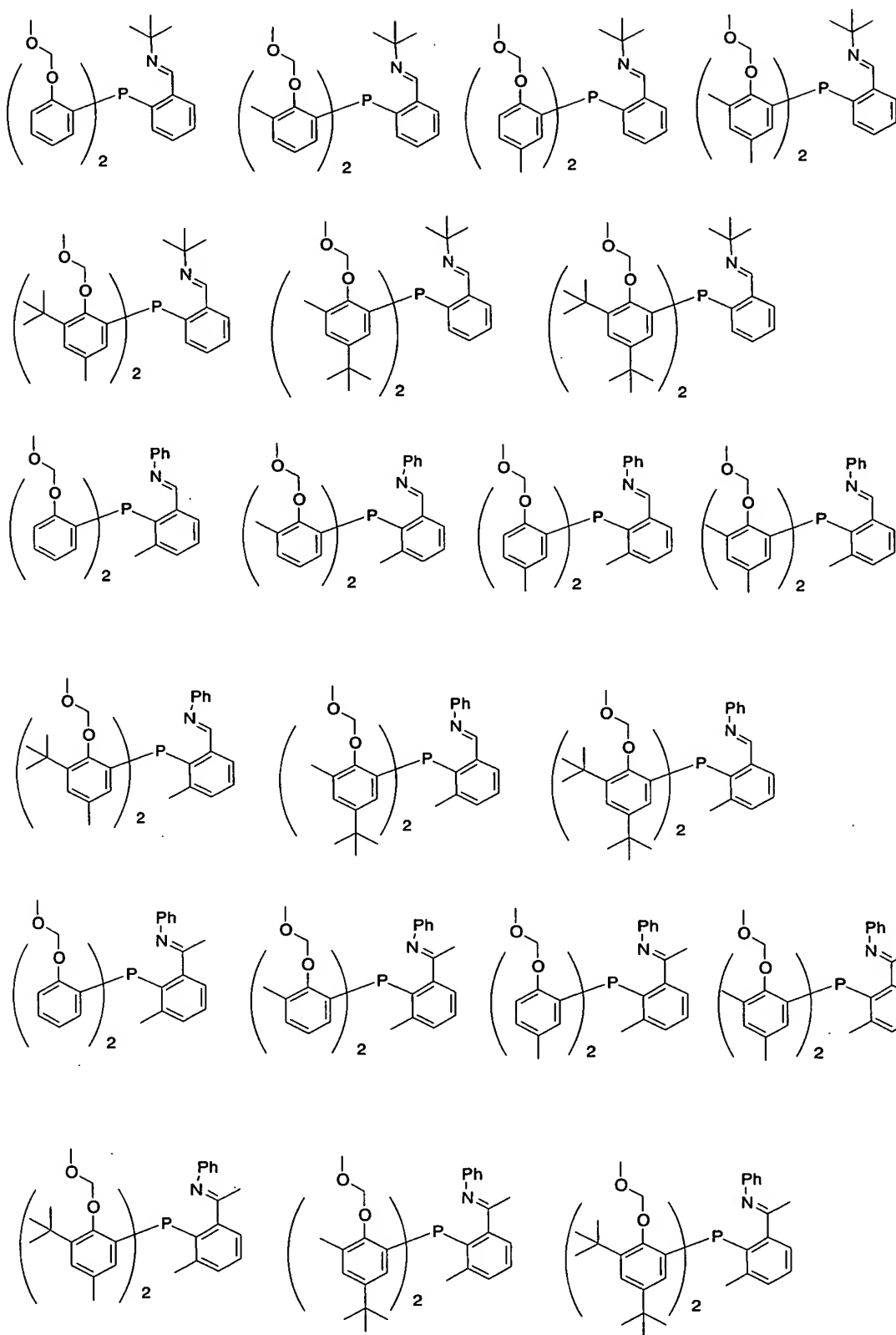


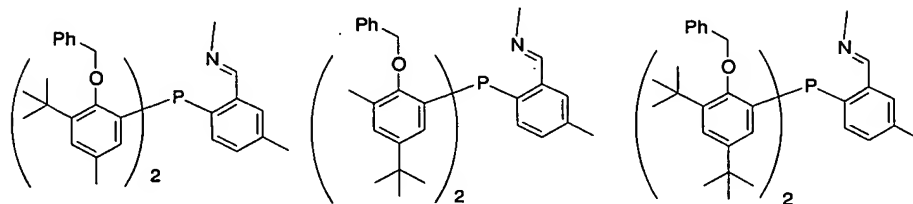
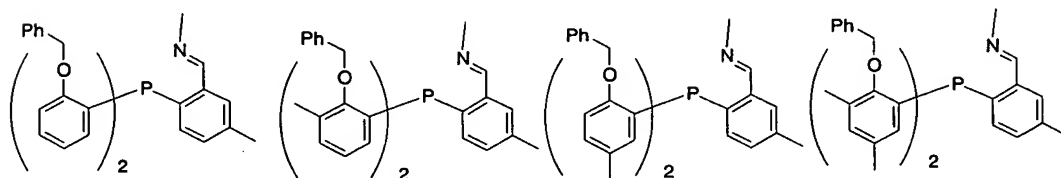
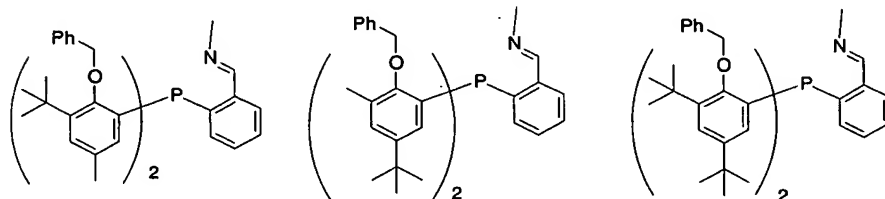
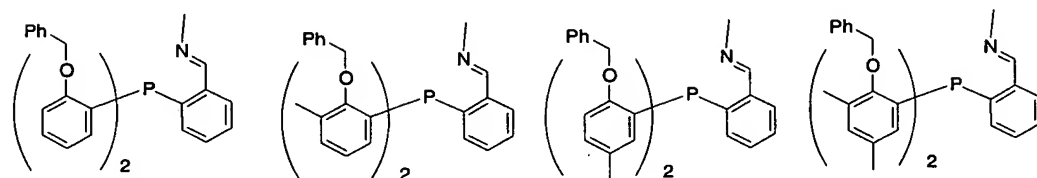
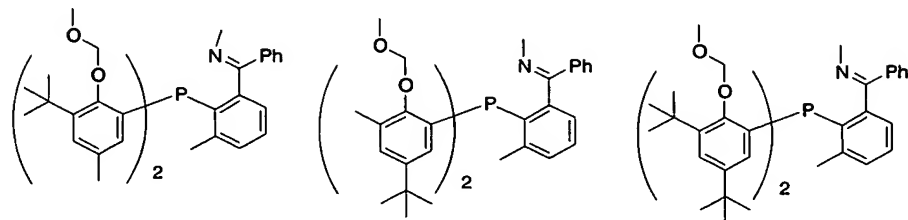
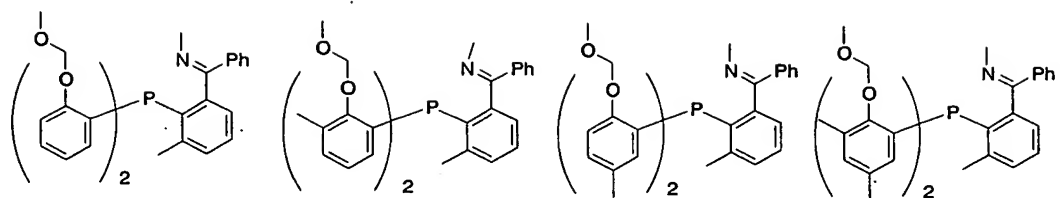
Specific examples of the phosphine compounds of formula (21B) include, for example, the following compounds:

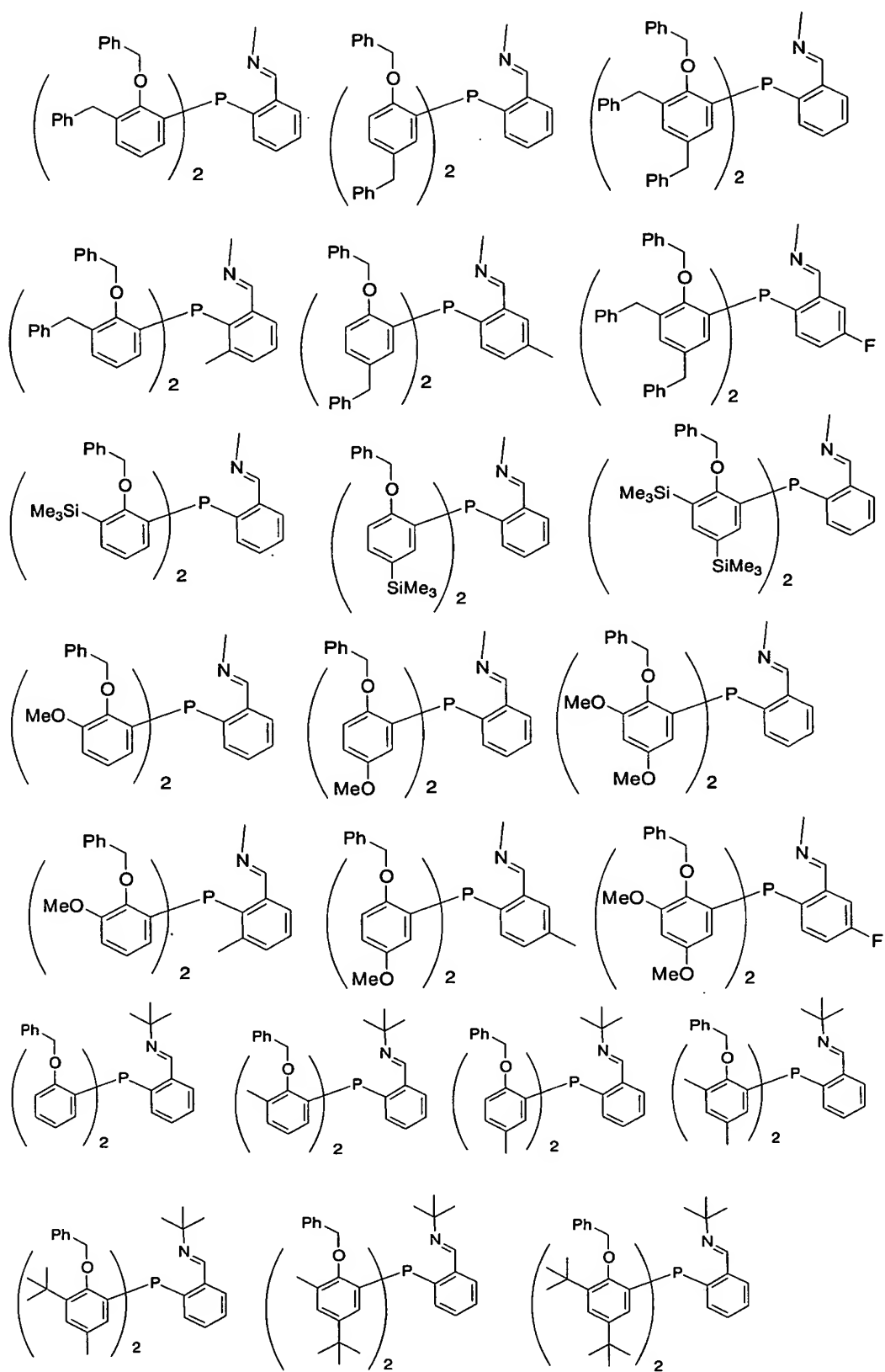


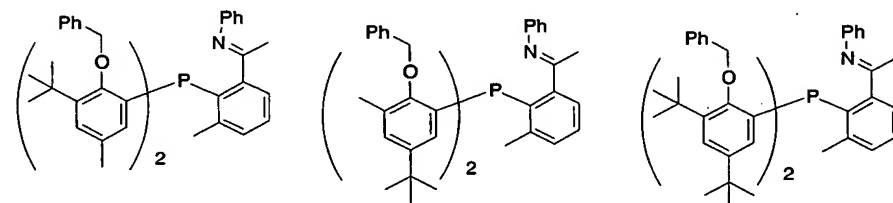
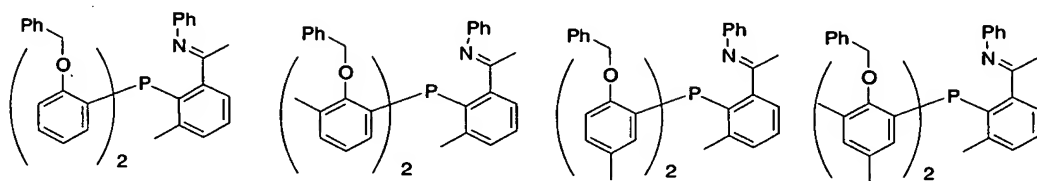
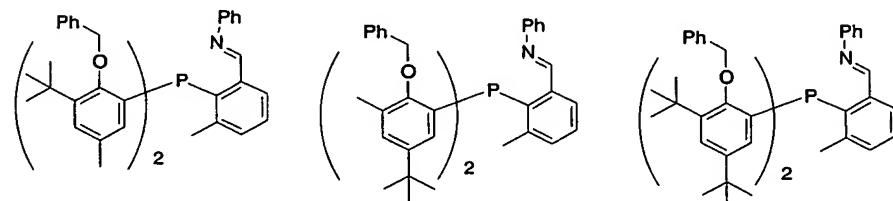
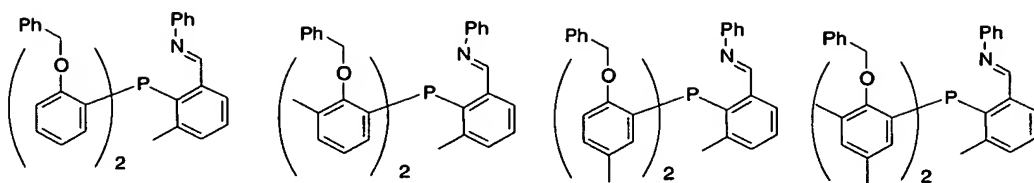


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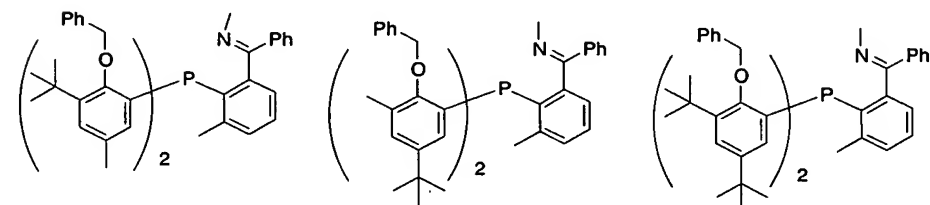
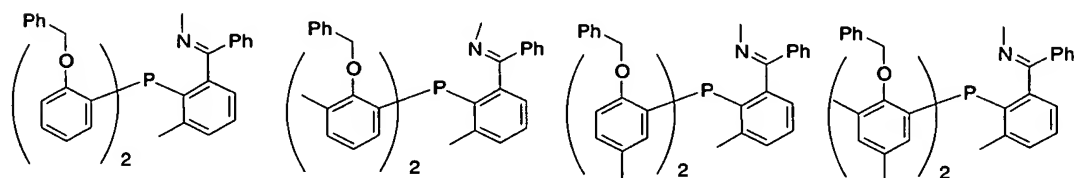


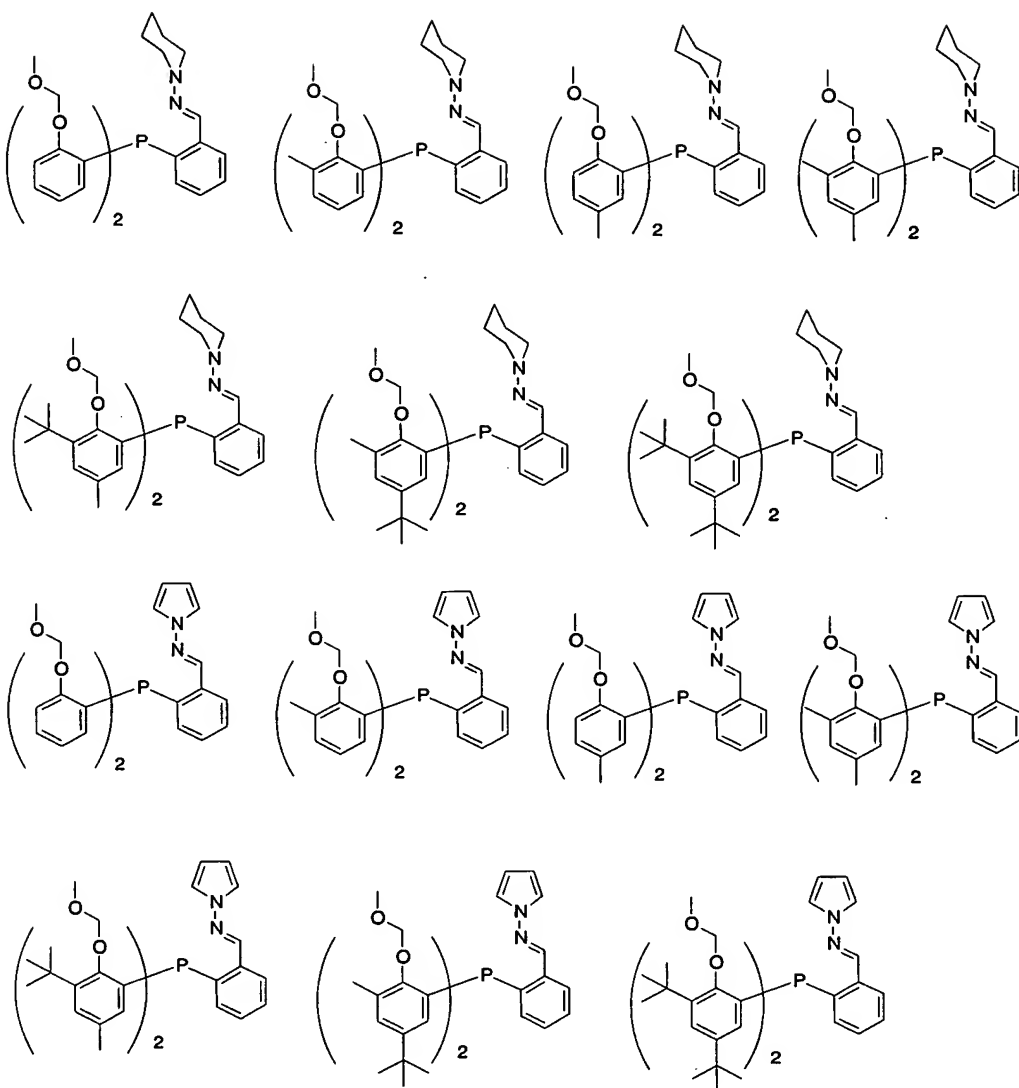






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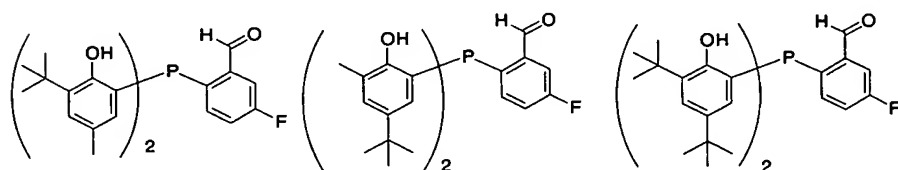
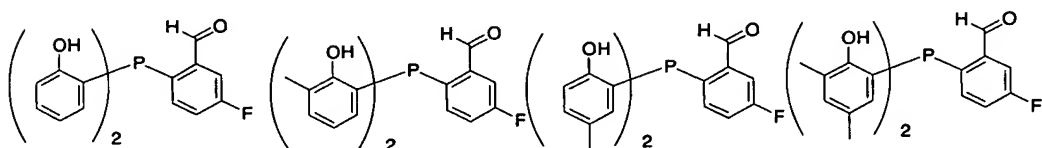
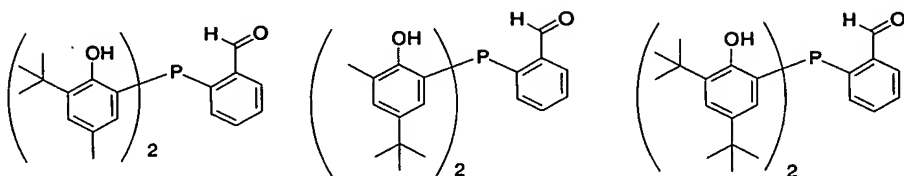
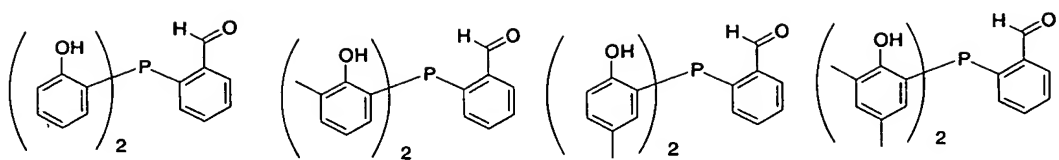
The compound of formula (21A), which corresponds to
 5 compounds of formula (1) wherein G^1 has the structure of G^{21} can
 be produced by reacting the phosphine carbonyl compound of
 formula (21C) with the organic compound of formula (21F).

While the reaction molar ratio between the phosphine
 carbonyl compound of formula 21C and the organic compound of
 10 formula (21F) is not particularly restricted, the ratio is
 preferably in the range of 1:0.1 to 1:10, more preferably in

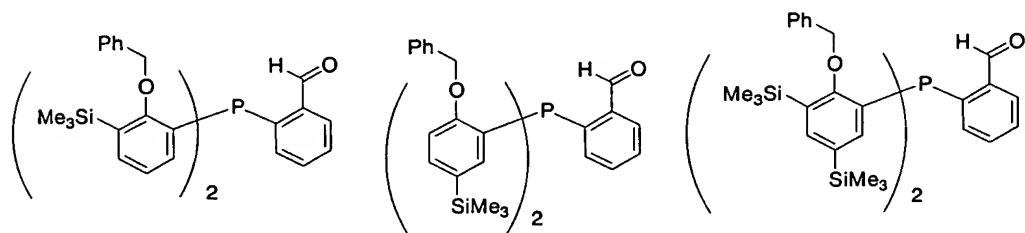
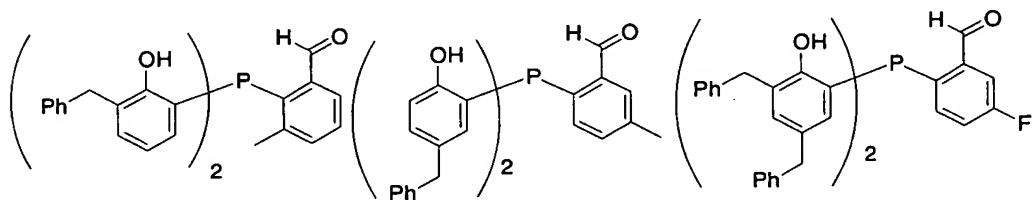
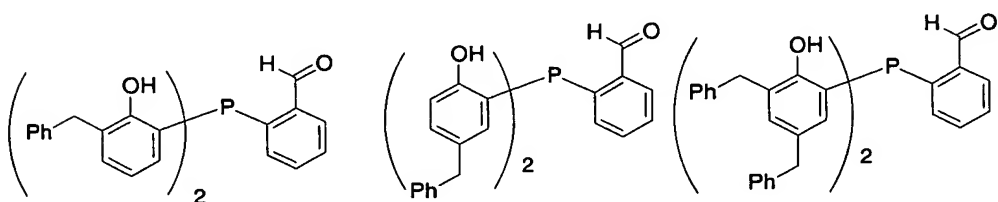
the range of 1:0.5 to 1:5.

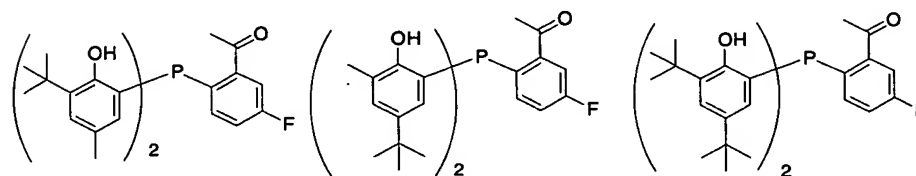
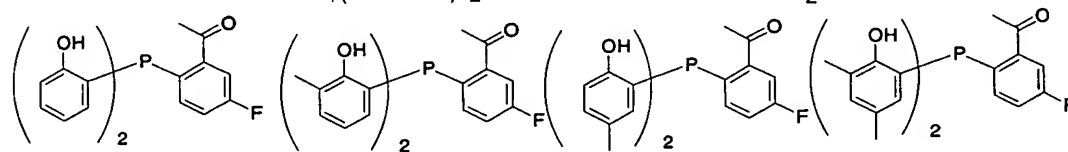
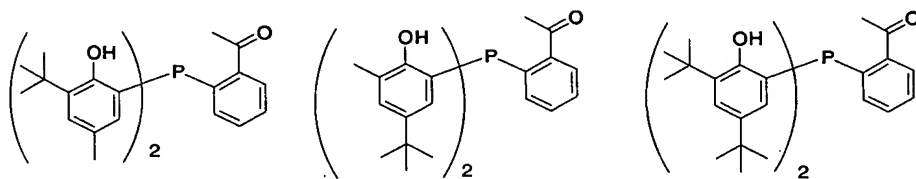
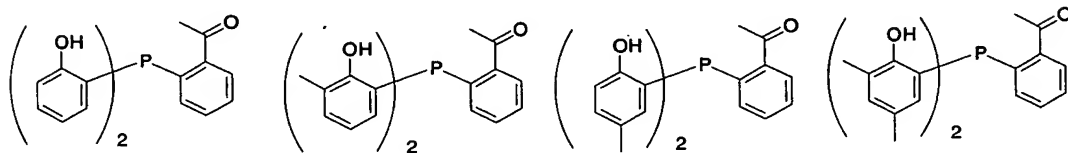
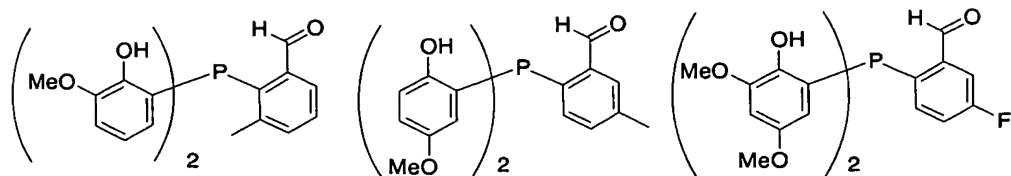
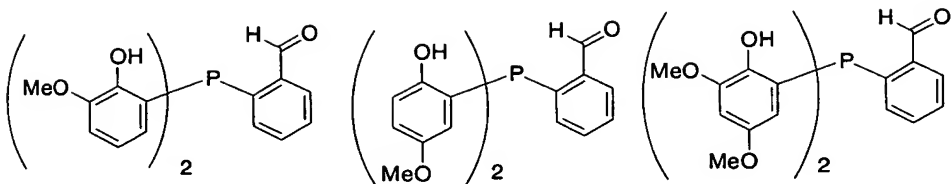
The reaction is usually performed in an inert solvent. Examples of these solvents include, for example, aprotic solvents including aromatic hydrocarbon solvents such as benzene or toluene; aliphatic hydrocarbon solvents such as hexane or heptane; ether solvents such as diethyl ether, tetrahydrofuran or 1,4-dioxane; polar solvents such as acetonitrile, propionitrile, acetone, diethyl ketone, methyl isopropyl ketone, cyclohexanone or ethyl acetate; and hydrogenated solvents such as dichloromethane, dichloroethane, chlorobenzene or dichlorobenzene; and protonic solvents such as methanol, ethanol, isopropanol or butanol. These solvents may be used alone or as a mixture of at least two of them. The amount of these solvents is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the carbonyl compound of formula (7) .

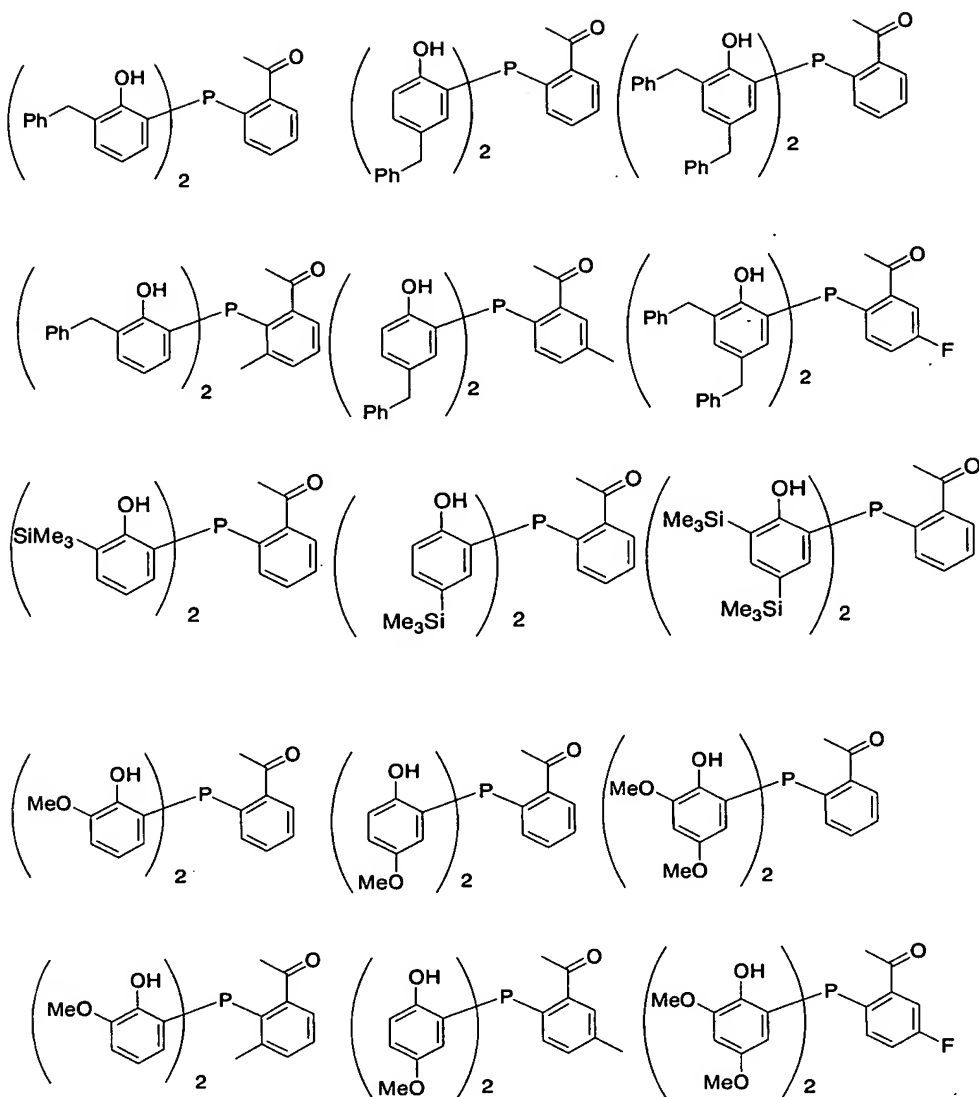
The reaction temperature is usually in the range of from -100°C or more to the boiling point or below of the solvent, preferably about -80 to 100°C. The phosphine compound of formula (21C), wherein G^1 in the formula denotes a hydrogen atom can be obtained from the reaction mixture by a conventional method, for example, by removing the solvent by evaporation. Specific examples of the phosphine carbonyl compound of formula (21C) include, for example, the following compounds:



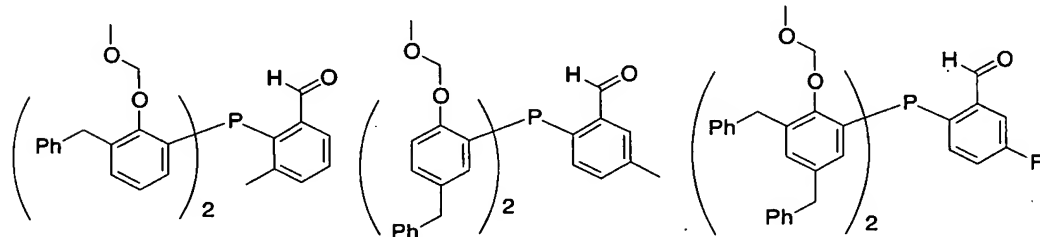
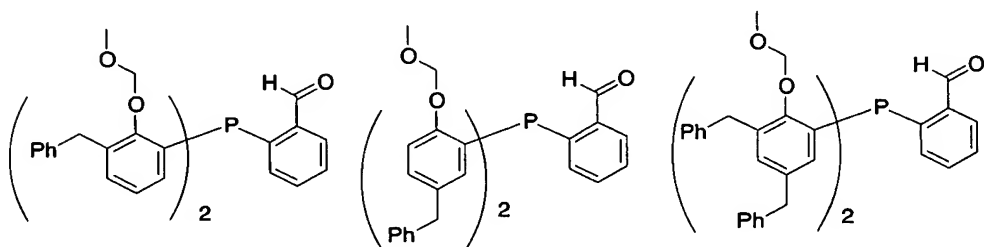
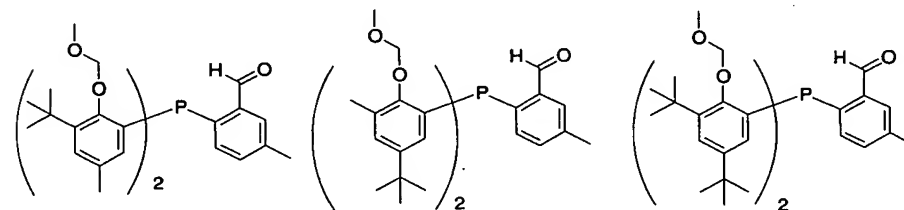
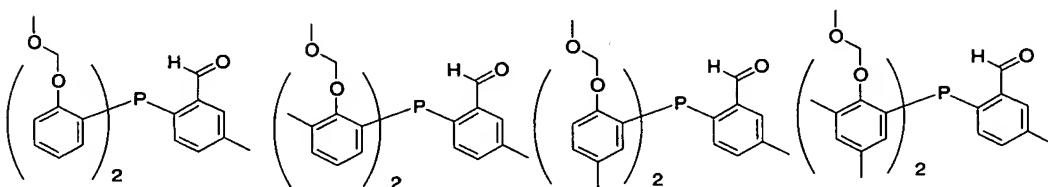
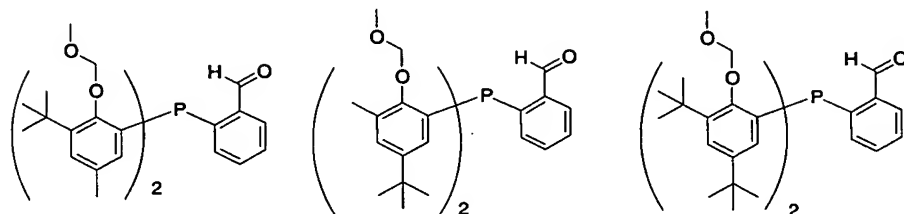
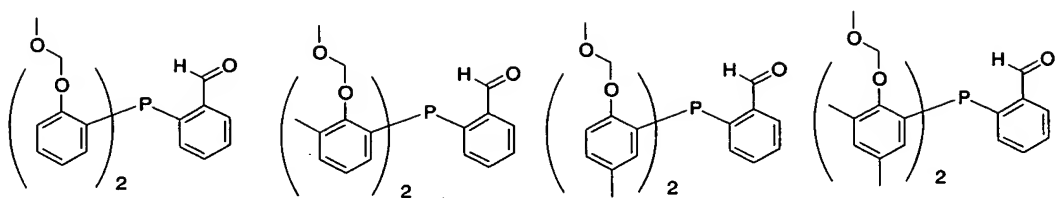
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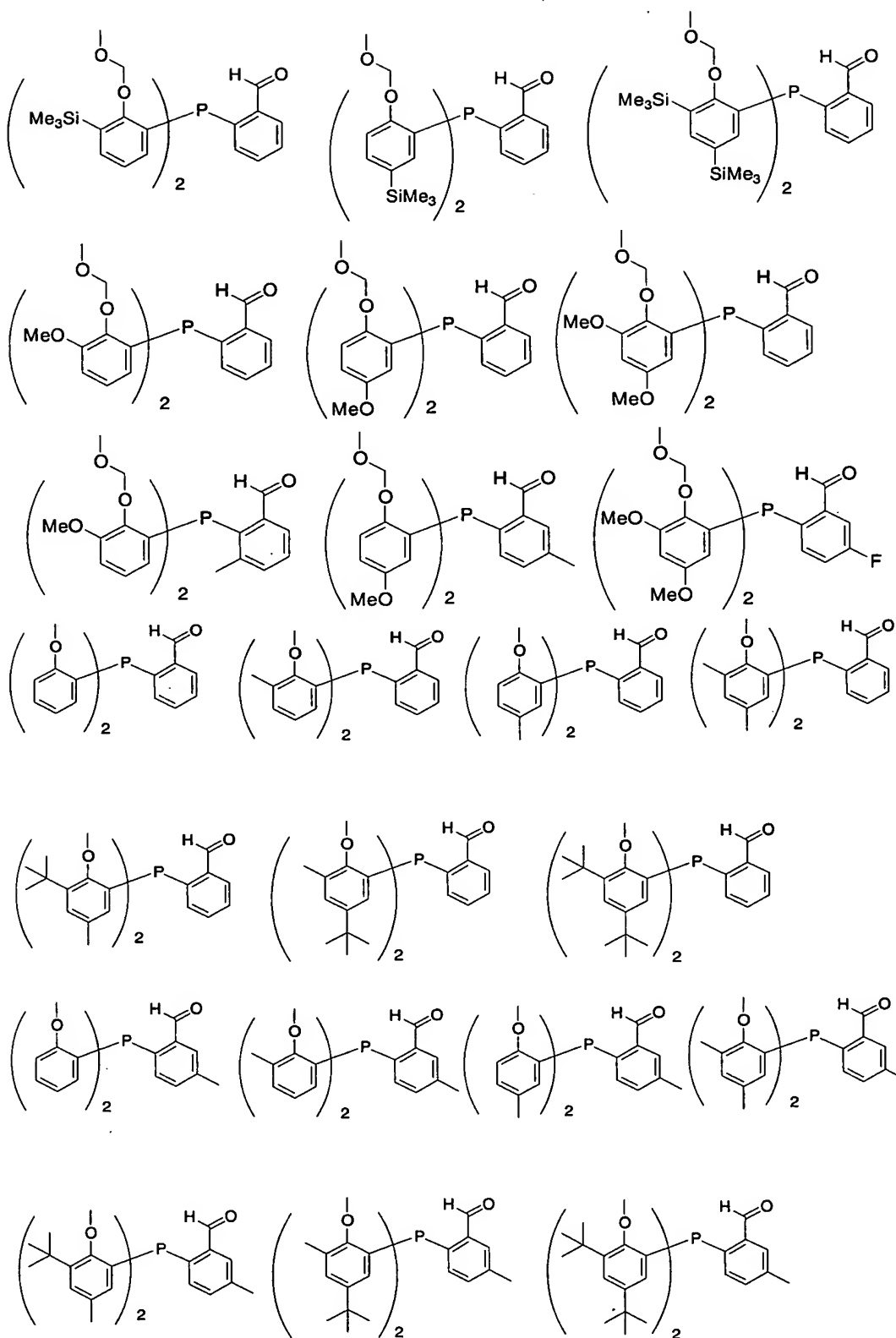


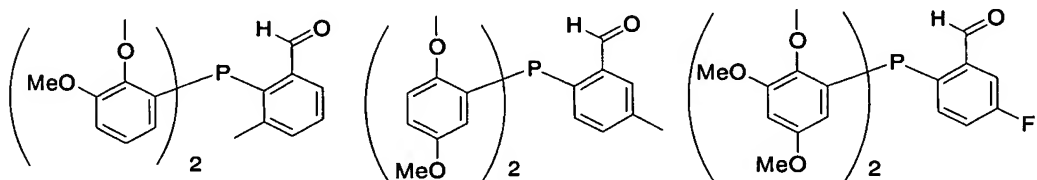
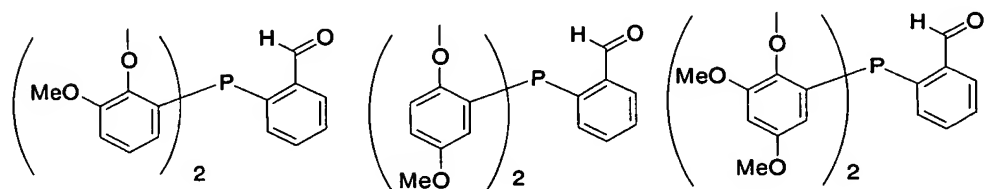
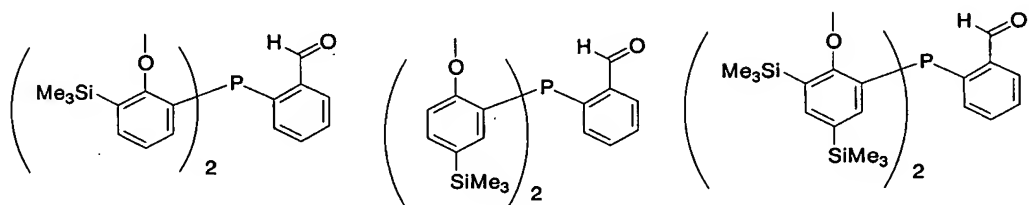
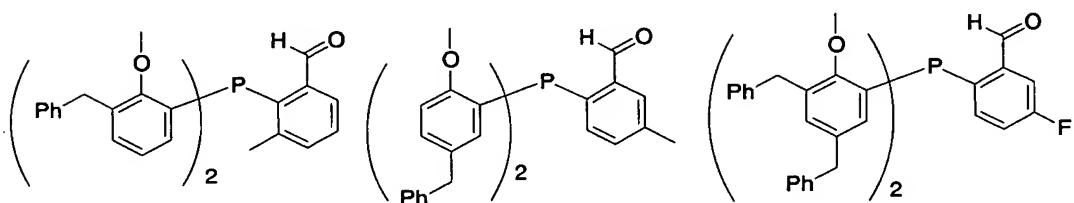
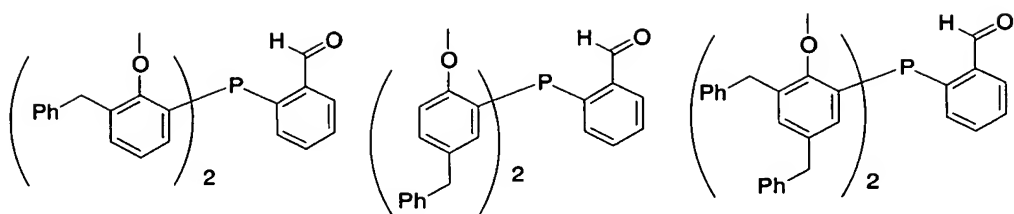


- 5 Specific examples of the organic compound of formula (21F) include methylamine, ethylamine, propylamine, butylamine, isobutylamine, tert-butylamine, pentylamine, hexylamine, aniline, 2-methylaniline, 4-methylaniline, 2,6-dimethylaniline, 2,6-diisopropylaniline,
- 10 pentafluoroaniline, aminopiperidine, aminopyrrole and the like. Specific examples of the carbonyl compound of formula (21C) include the following compounds:

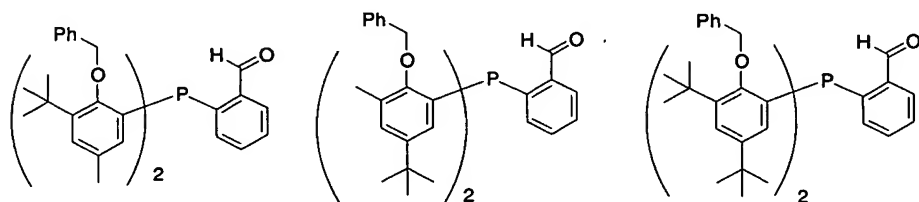
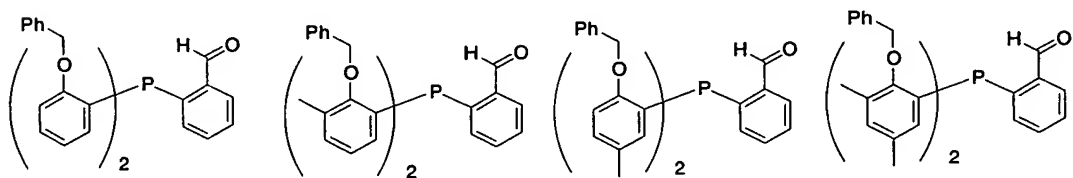


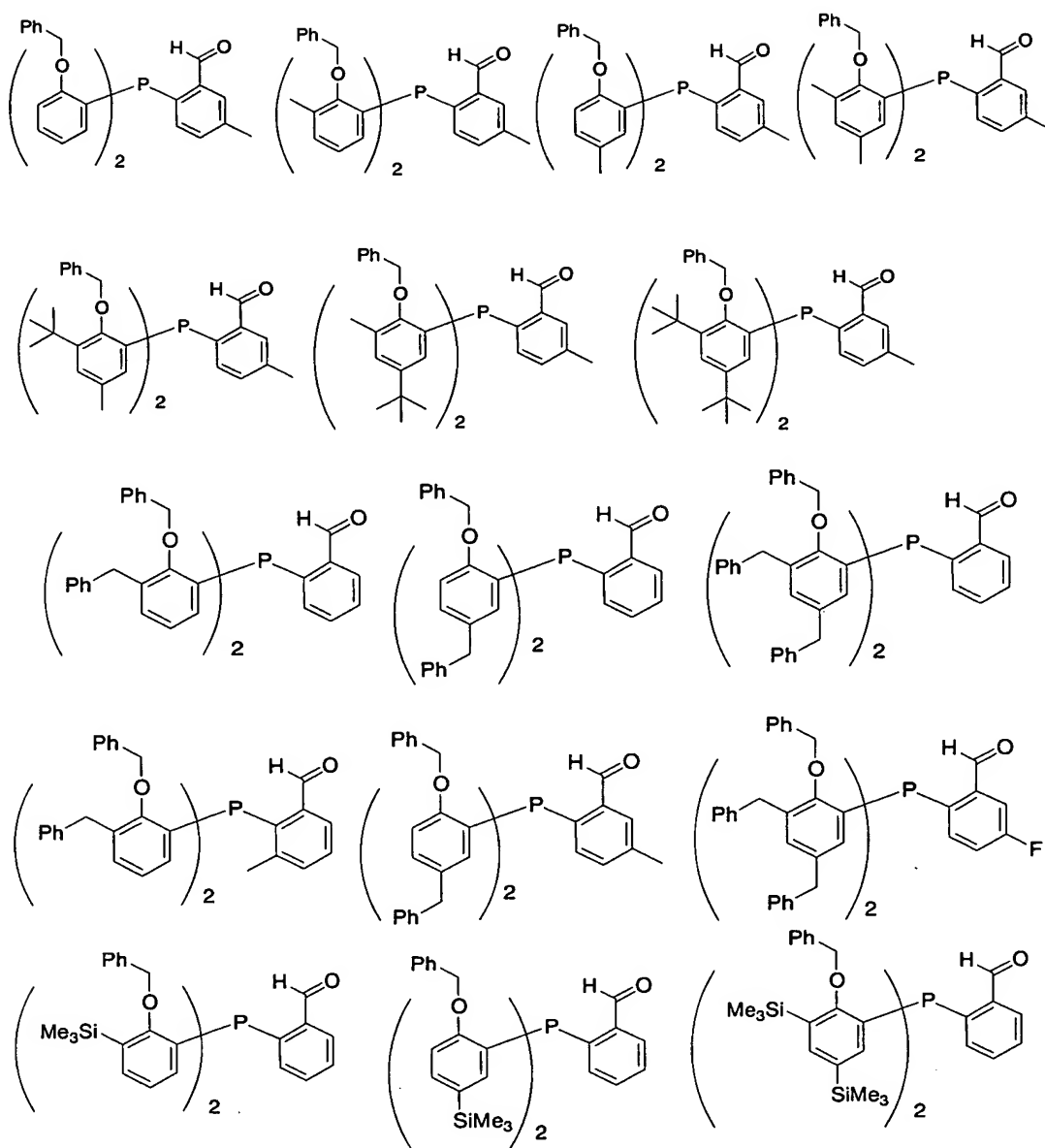
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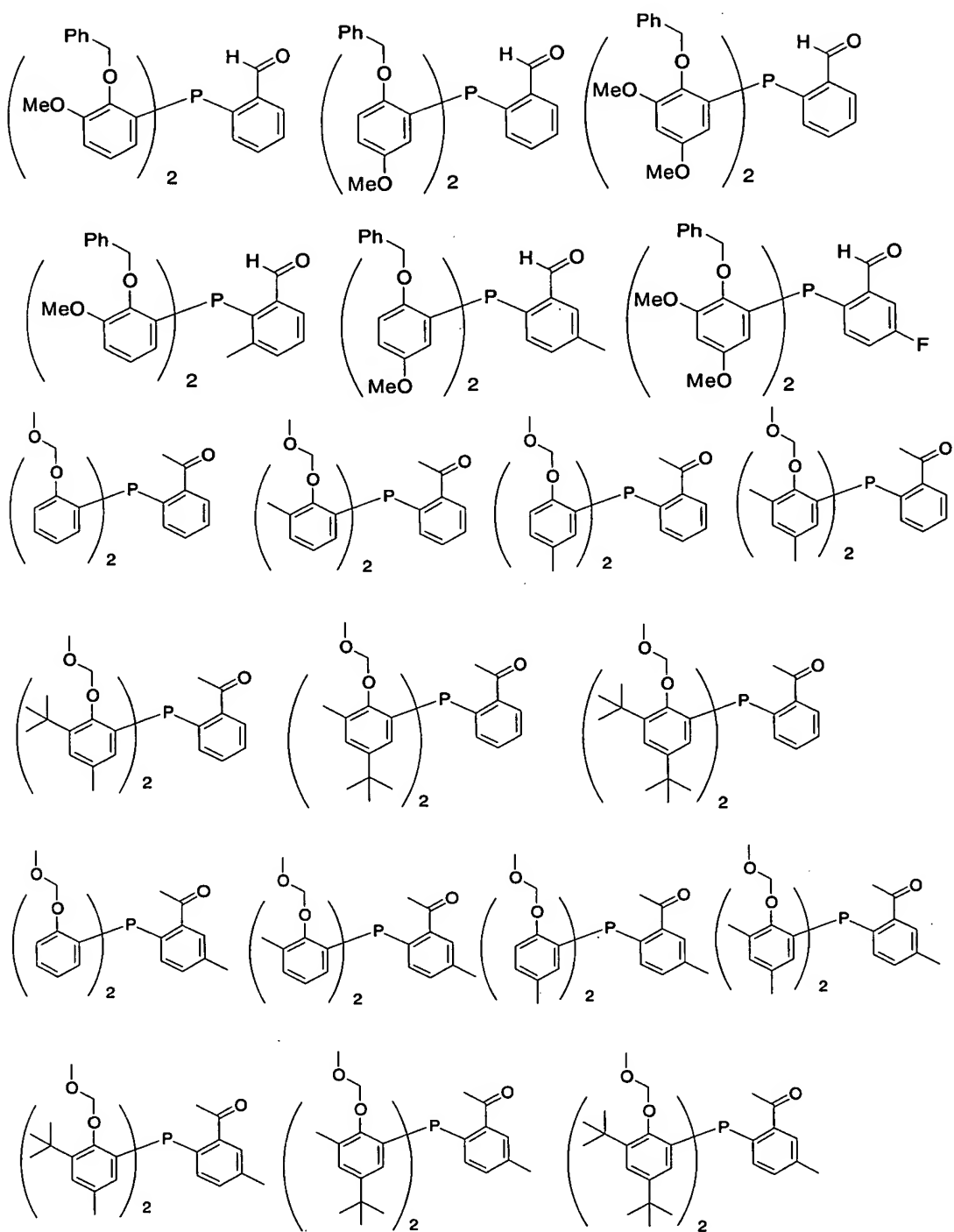


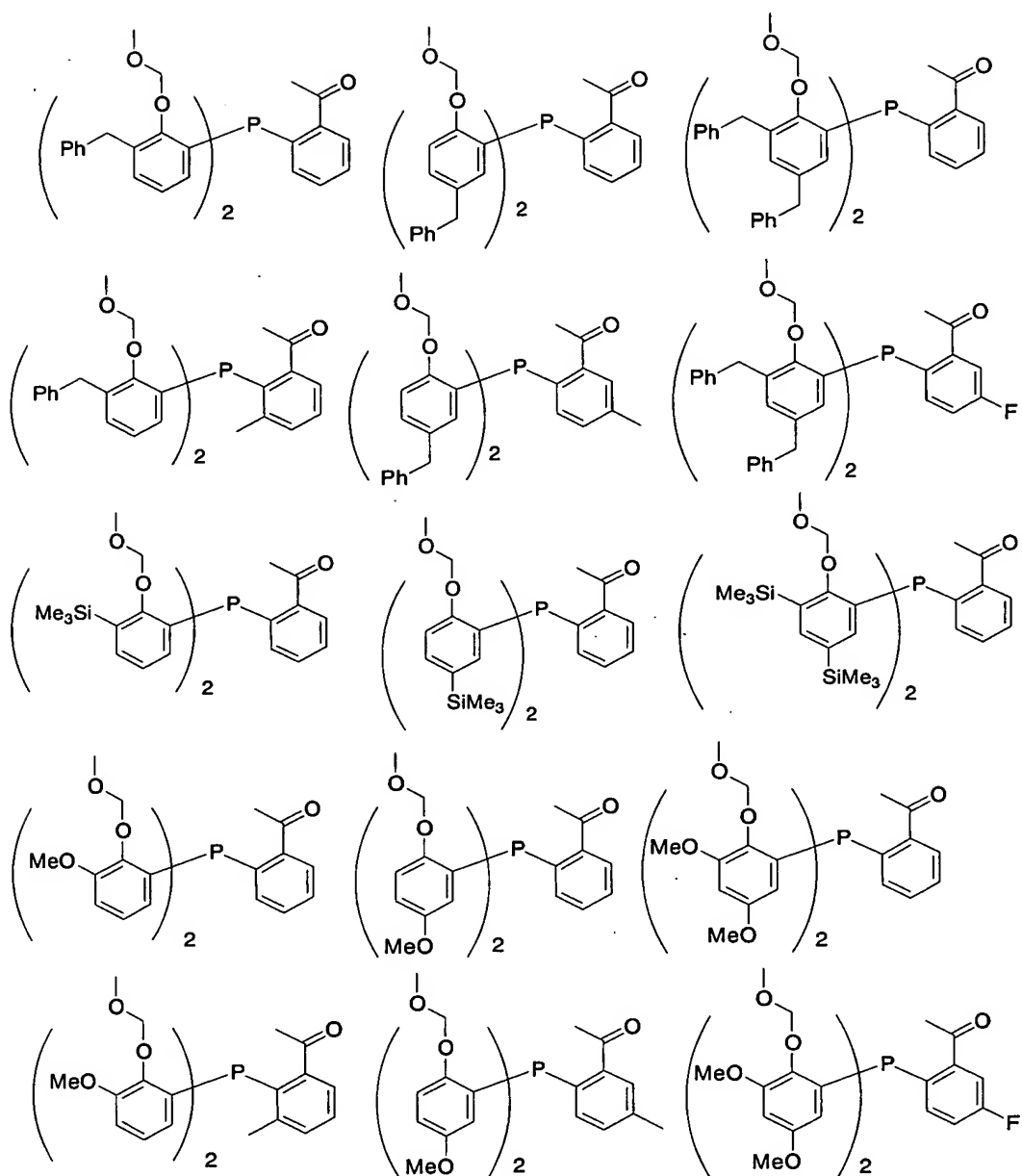
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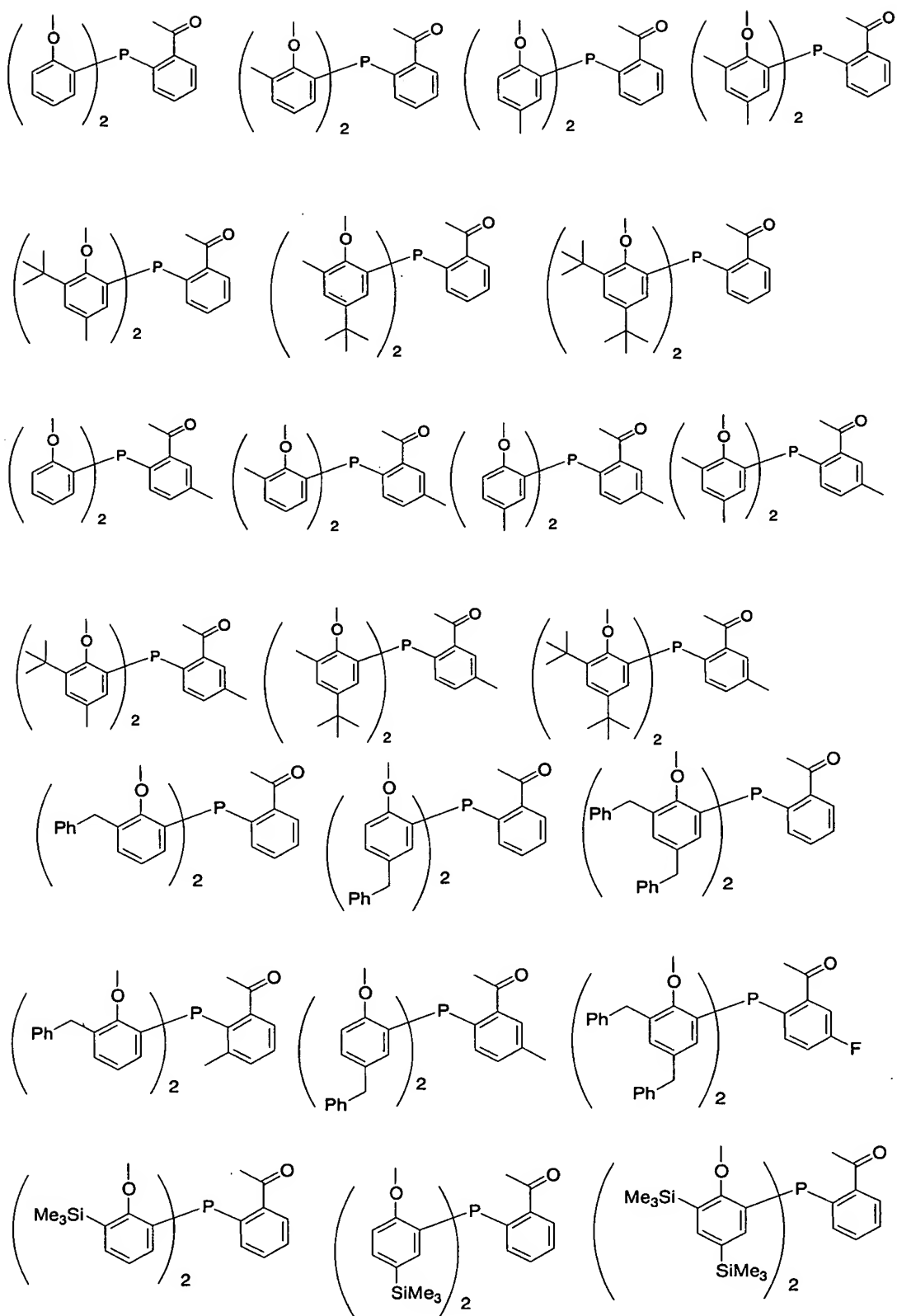


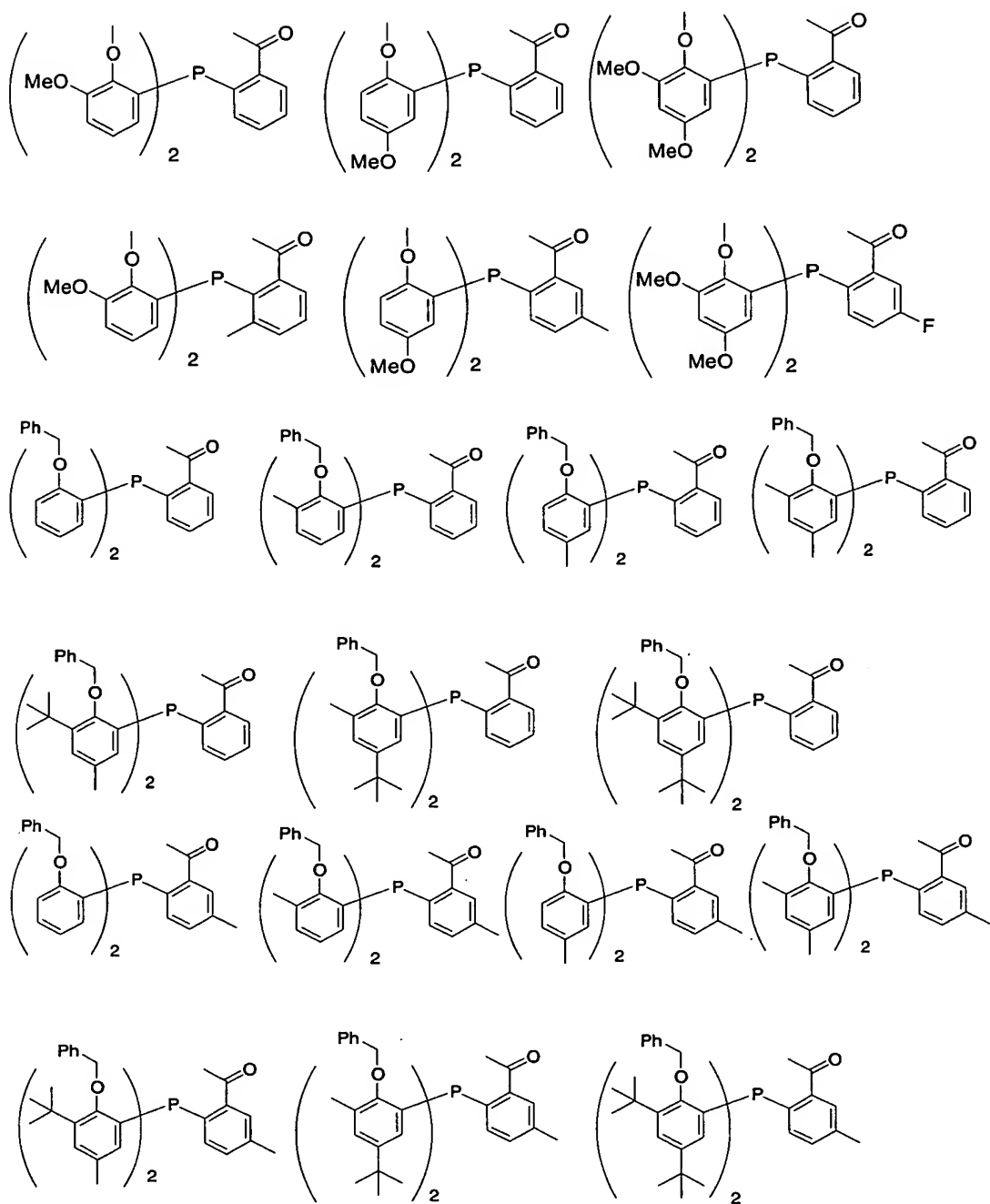


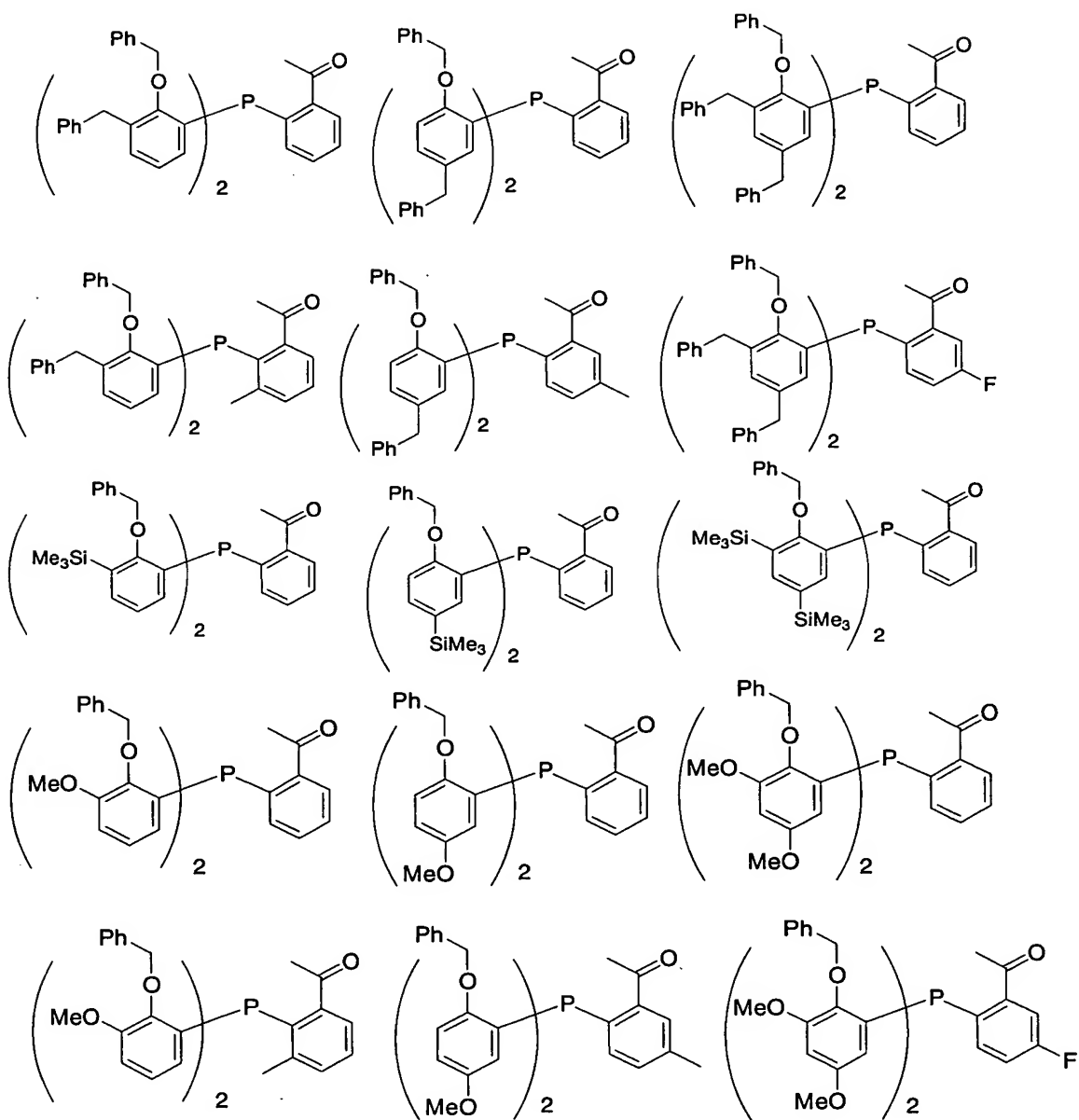
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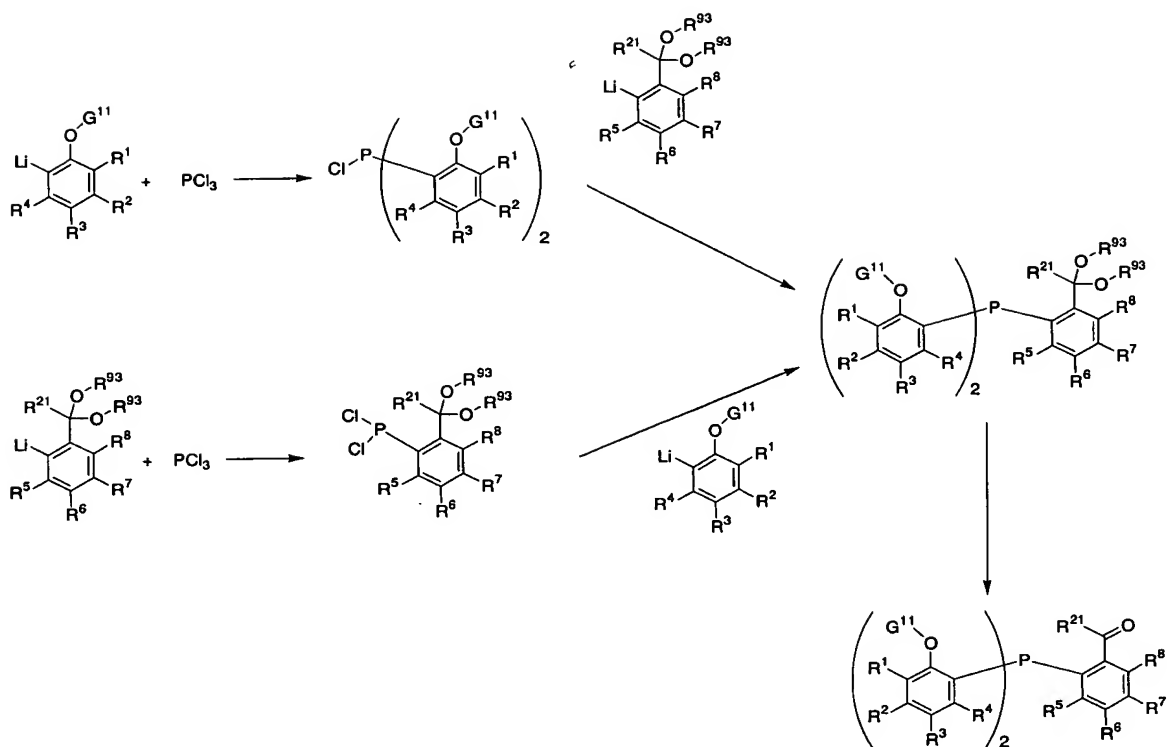








5 The phosphine compound of formula (21C) can be synthesized
by the following reaction route:

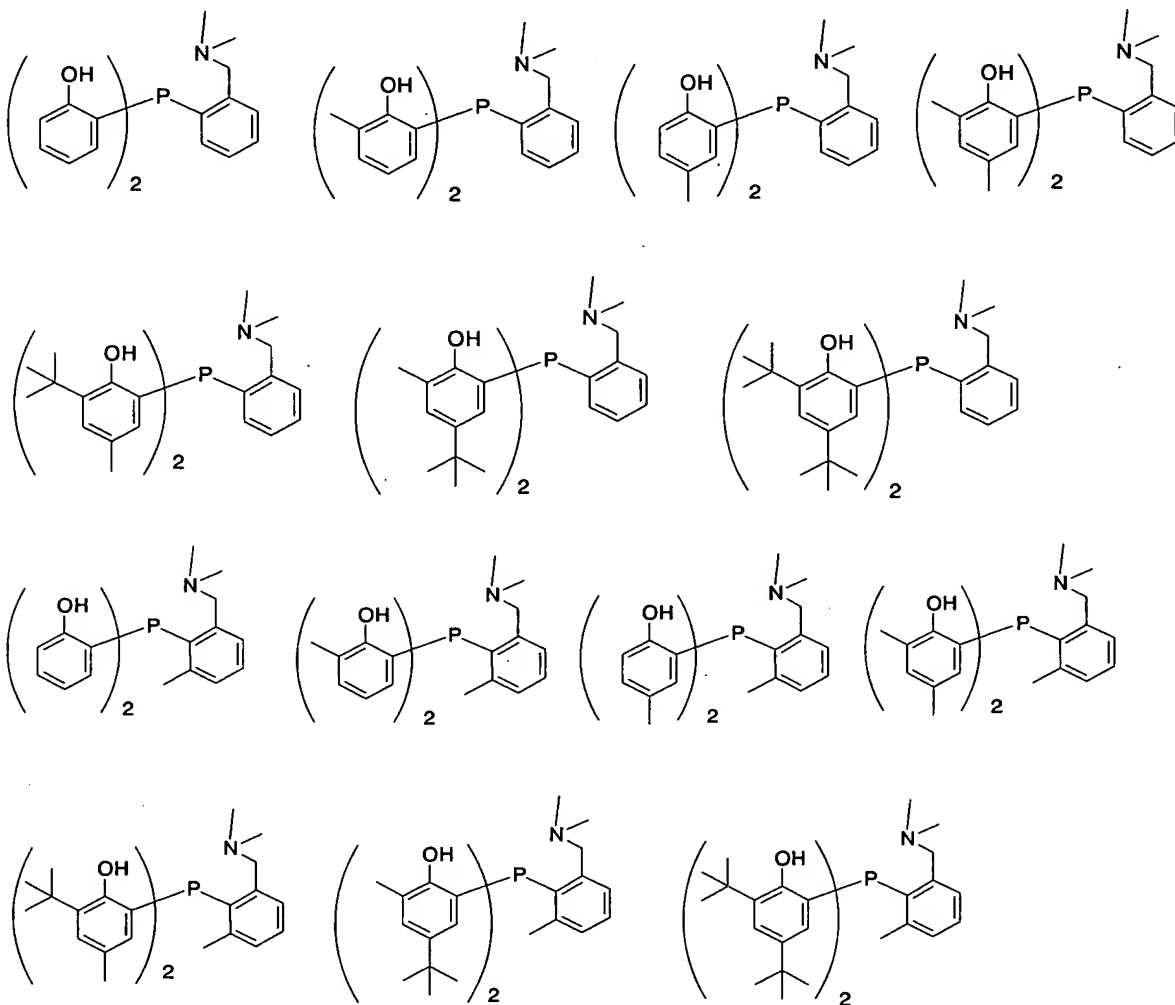


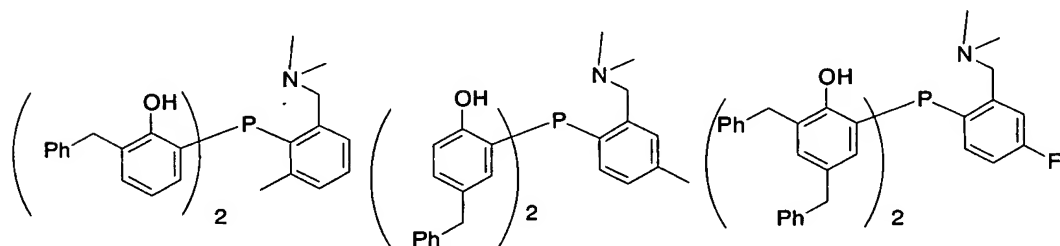
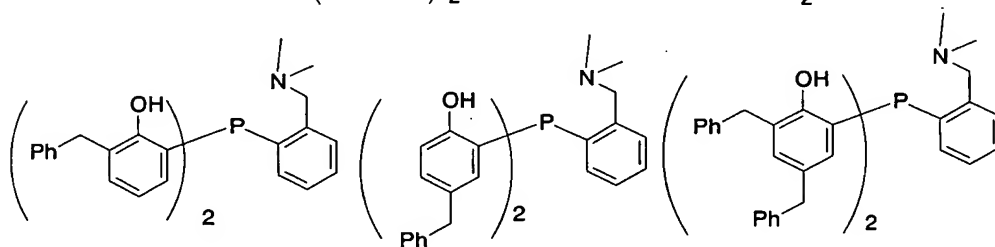
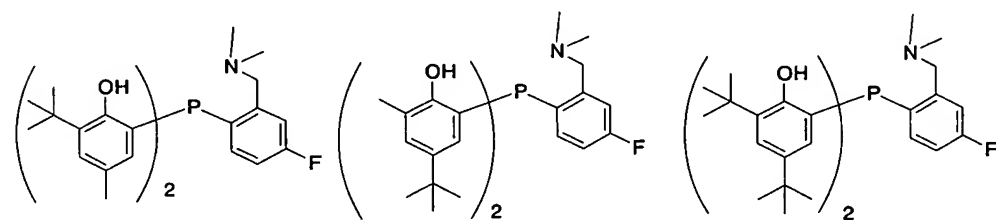
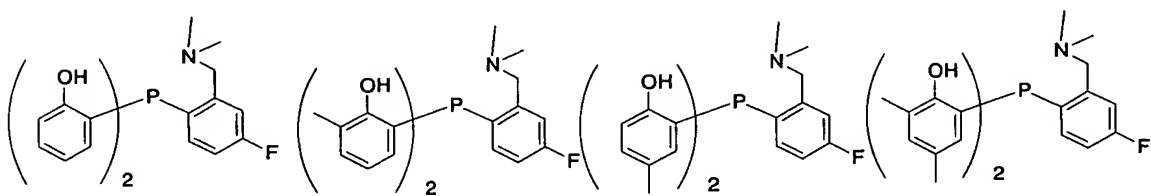
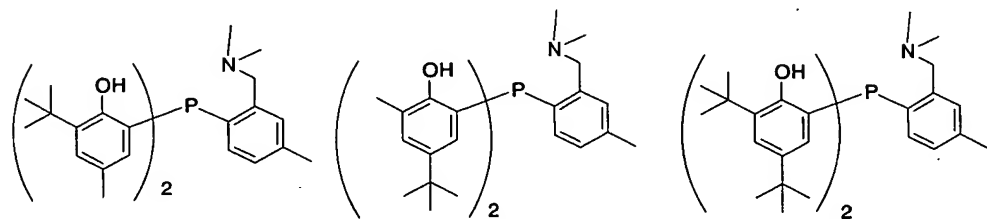
In the scheme, $R^1, R^2, R^3, R^4, R^5, R^6, R^7, R^8, R^{21}$ and G^{11} have the same meanings as described above, and R^{93} denotes a substituted or unsubstituted alkyl or aralkyl group, or a cyclic alkylene group by linking together.

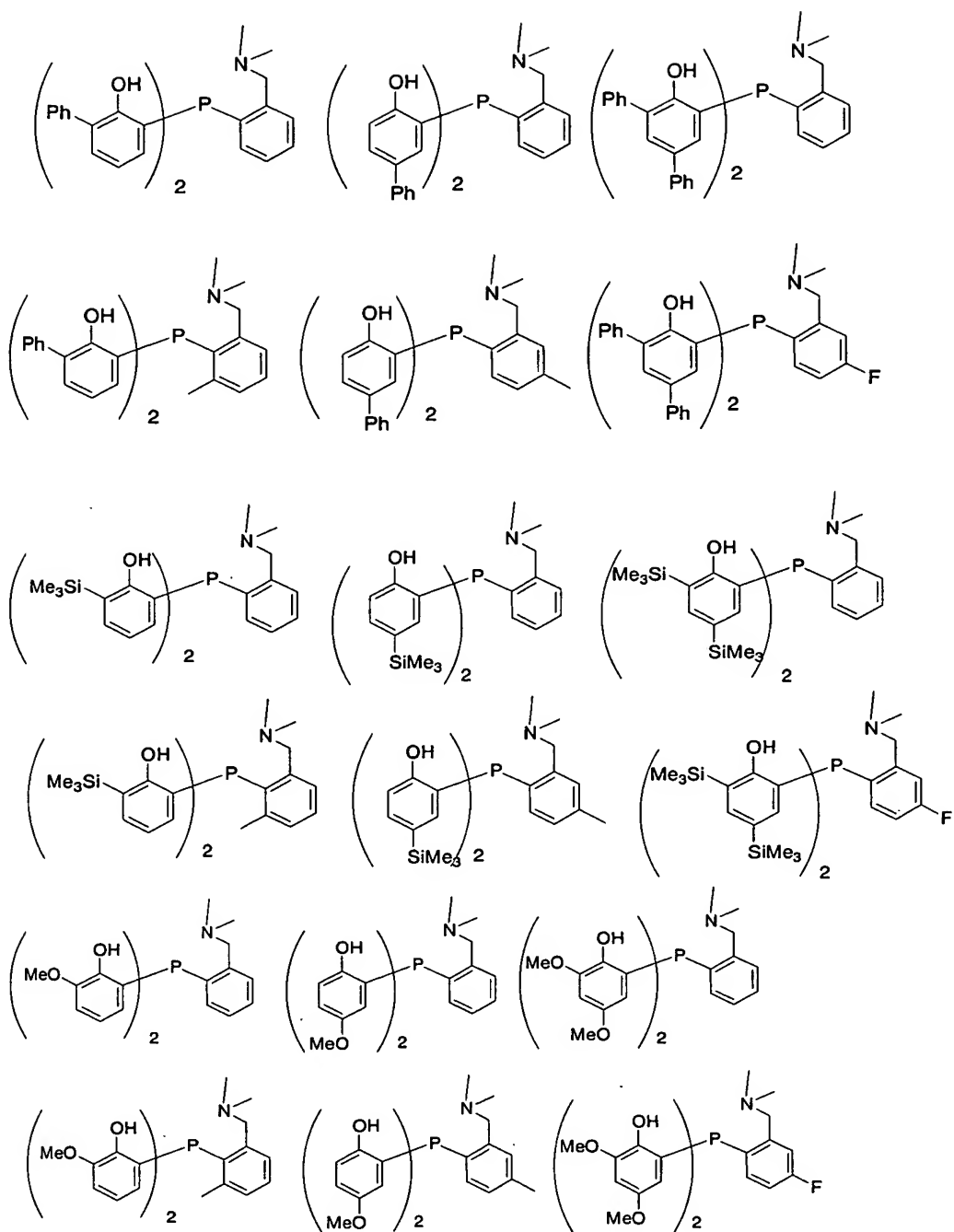
- 5 In the compound of formula (22A), which corresponds to the compound of formula (1), wherein G^1 has the structure of G^{22} , and A^1 is P, can be produced by deprotecting the compound of formula (22B) wherein A^1 is P, in a similar manner as the compound wherein A^1 is N as explained above. The compound of
- 10 formula (22B) wherein A^1 is P, can be synthesized from the compound of formula (22E) wherein which A^1 is P, in a similar manner as the compound wherein A^1 is N. The compound of formula (22E) wherein A^1 is P, can be synthesized from a phosphine-substituted halogenated aryl as a precursor, for example, by reacting
- 15 1-(α -chloromethyl)-2-bromobenzene with a mono-substituted

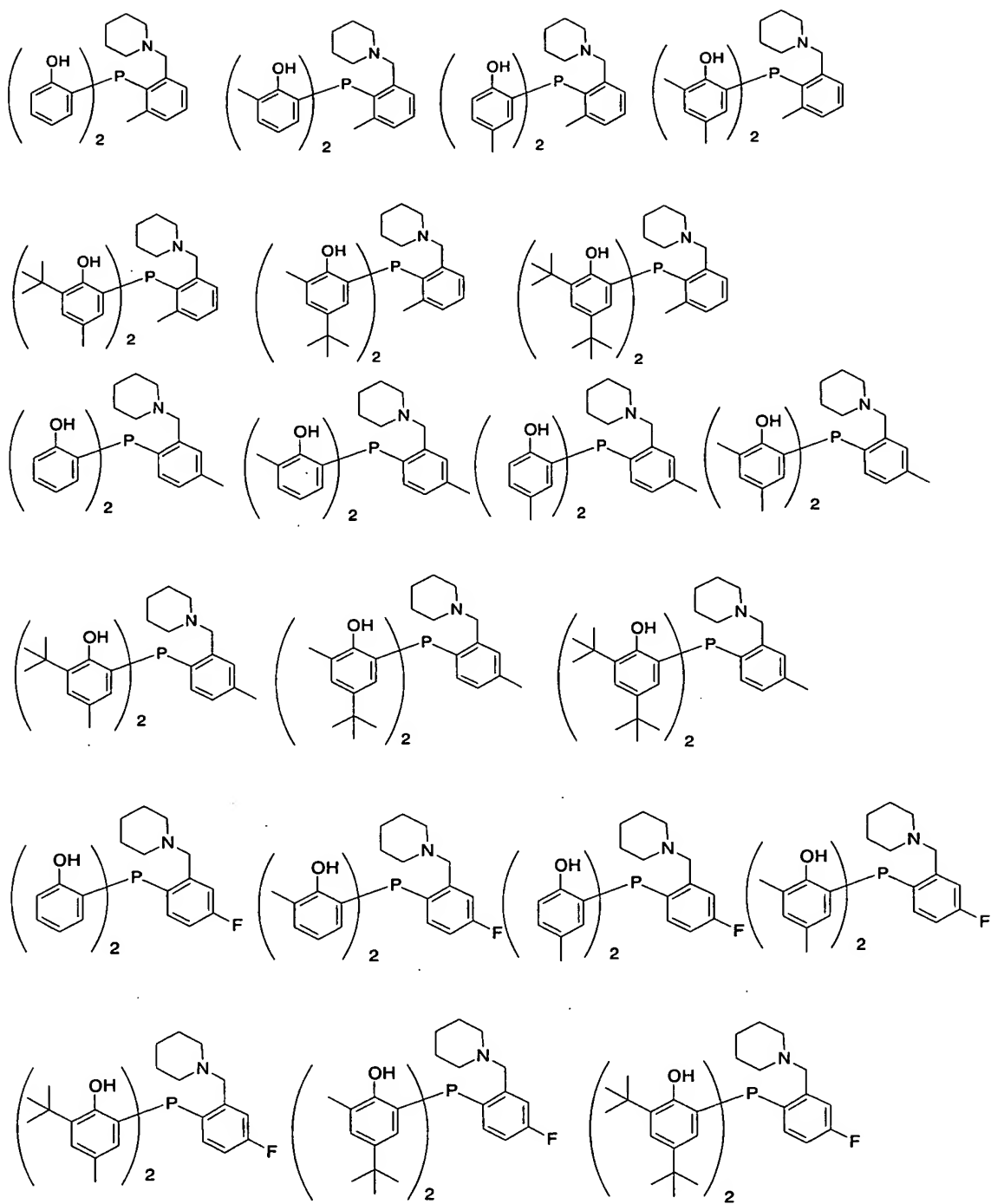
phosphine in the presence of a base as described in Journal of Praktische Chemie (Leipzig), vol. 330, p674, 1988.

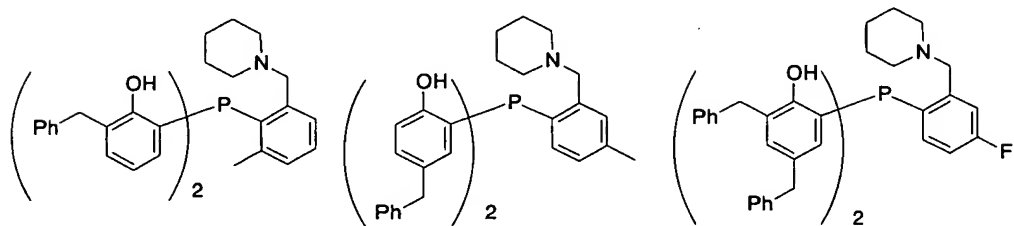
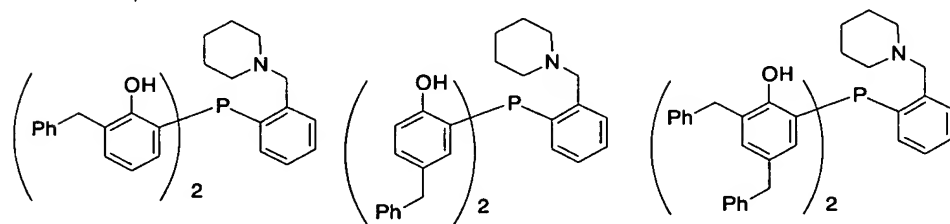
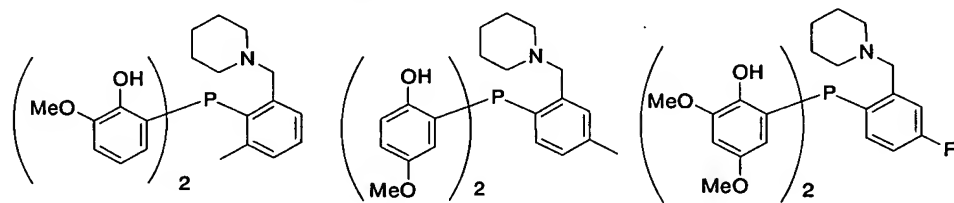
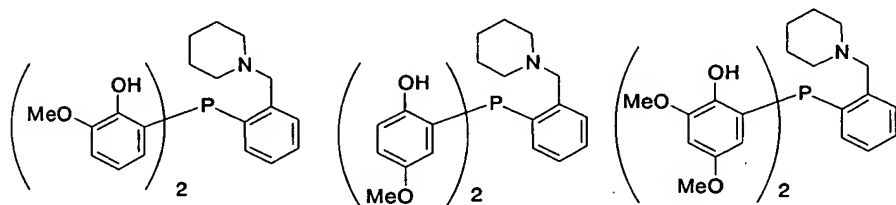
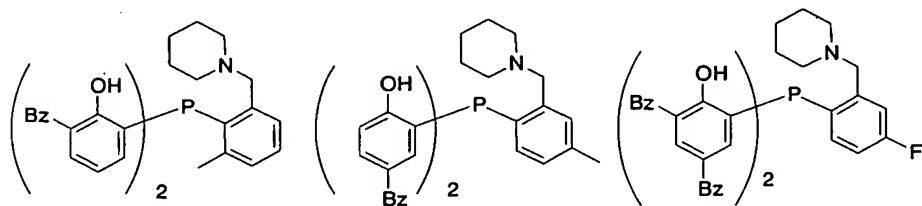
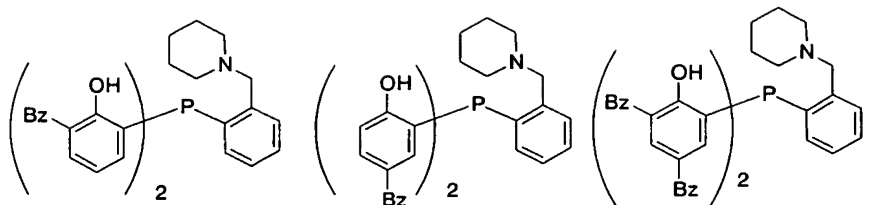
Specific examples of the compound of formula (22A), which corresponds to the compounds of formula (1) wherein G^1 has the structure of G^{22} include, for example, the following compounds:



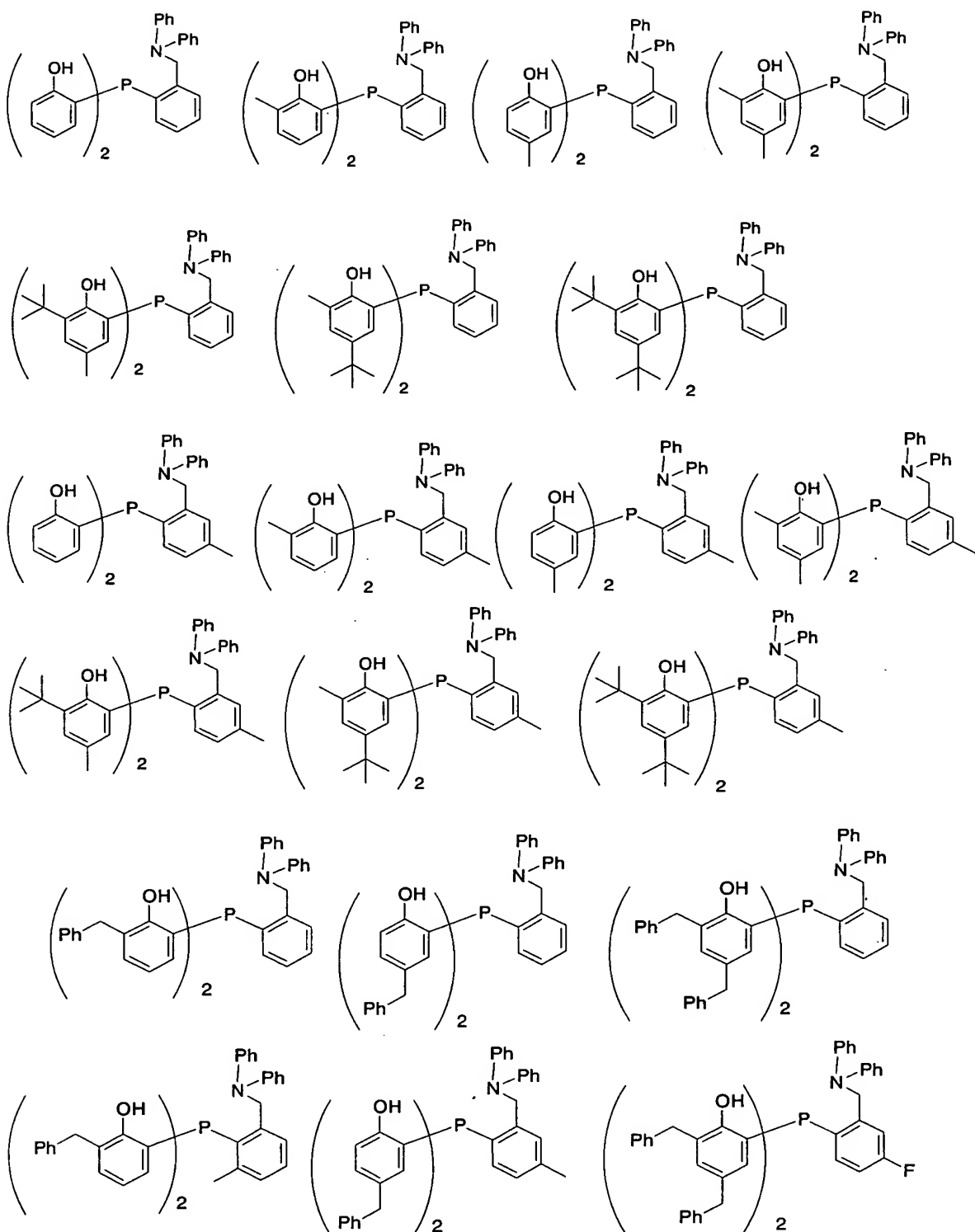


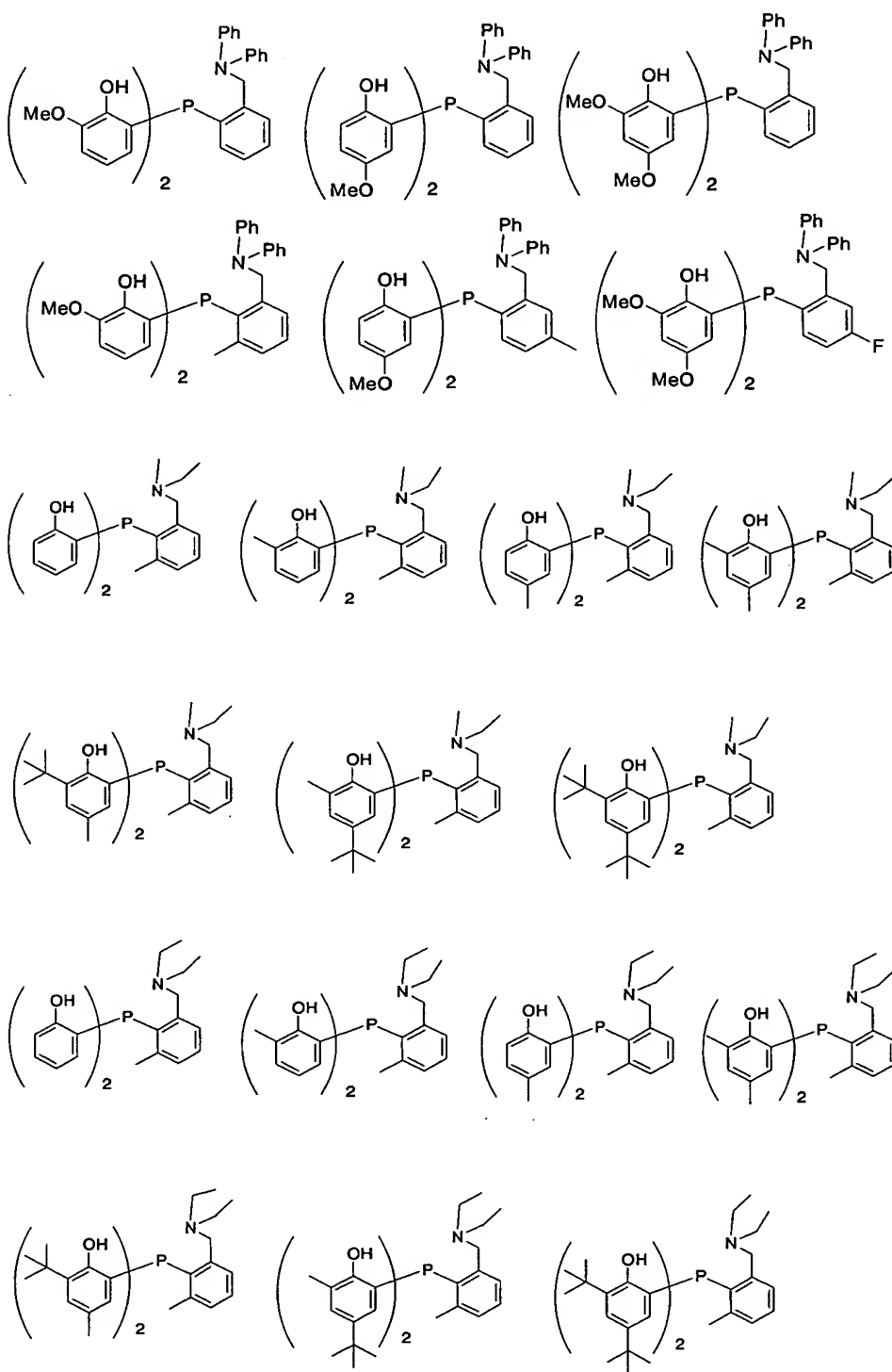


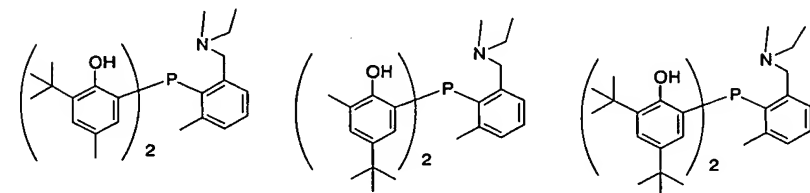
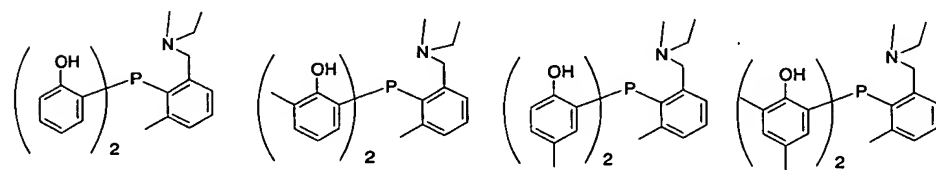
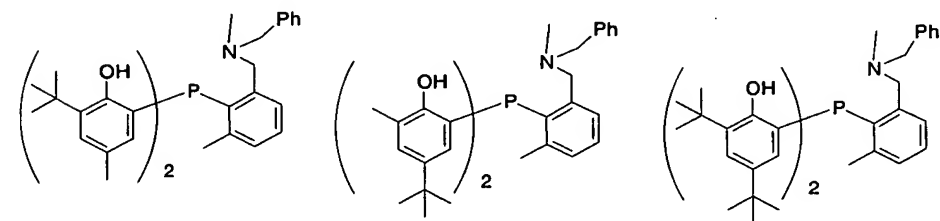
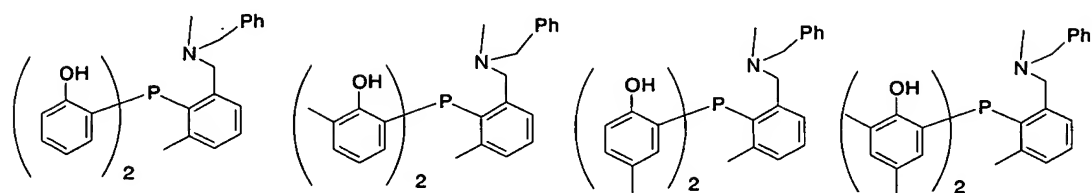
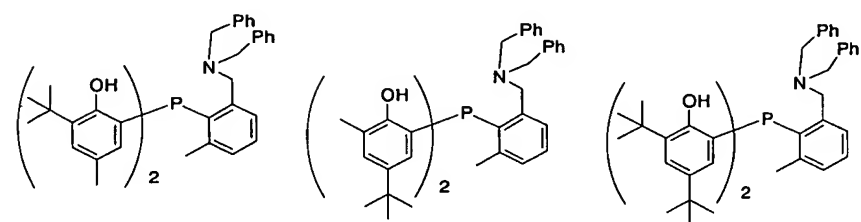
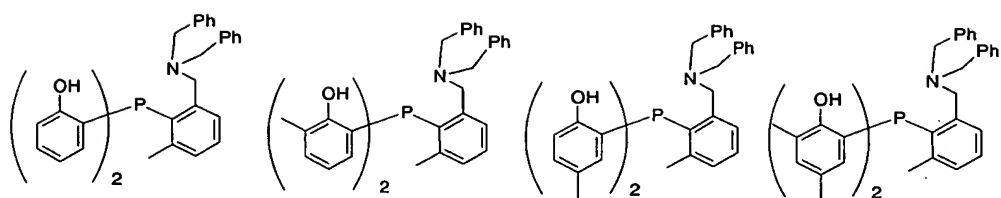


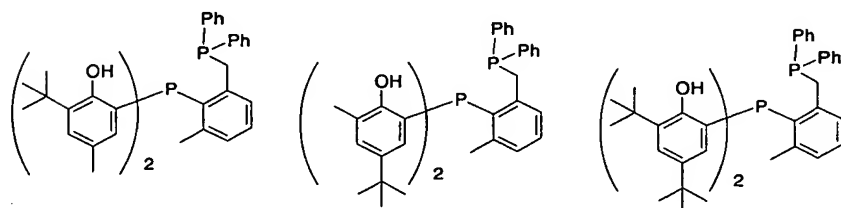
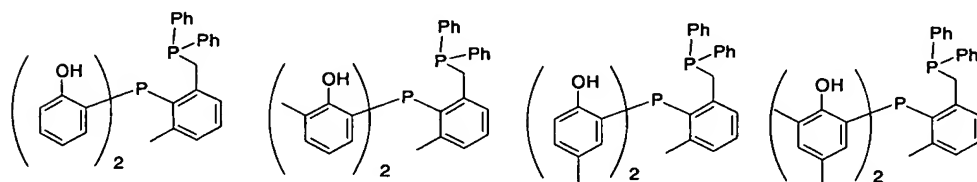
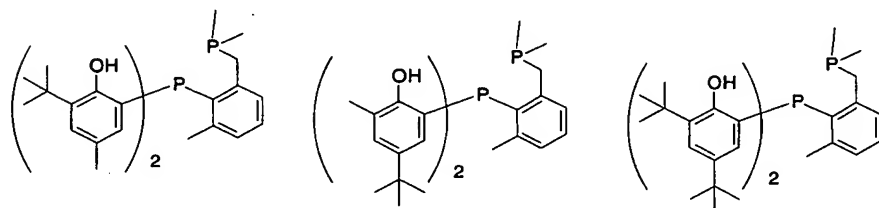
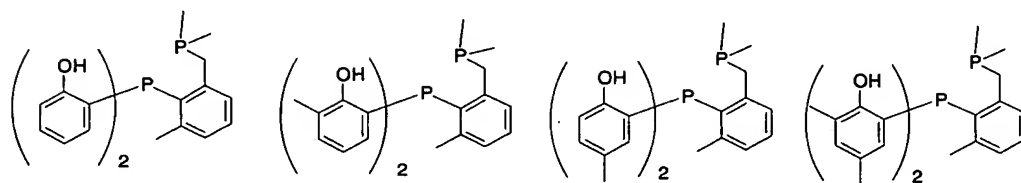
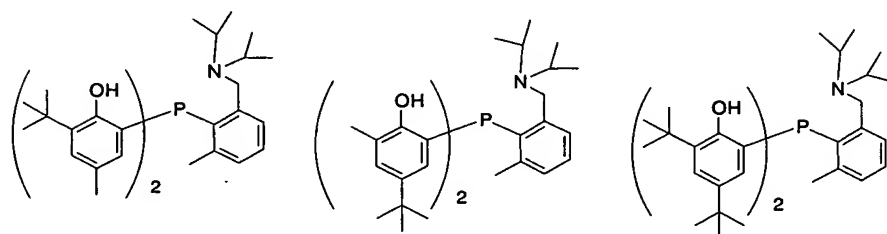
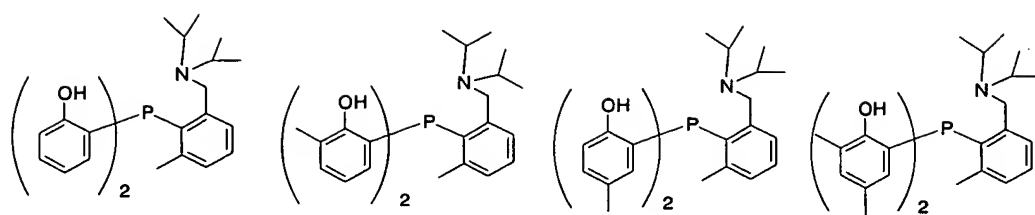


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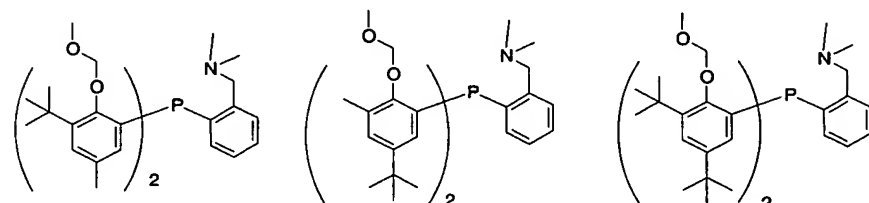
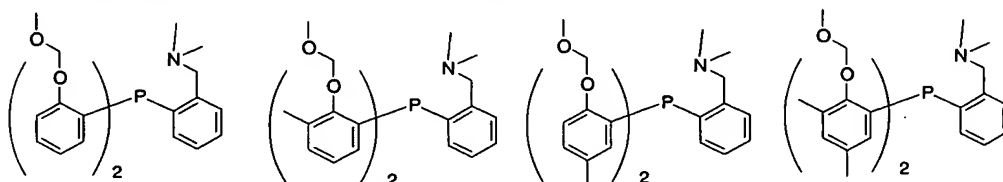




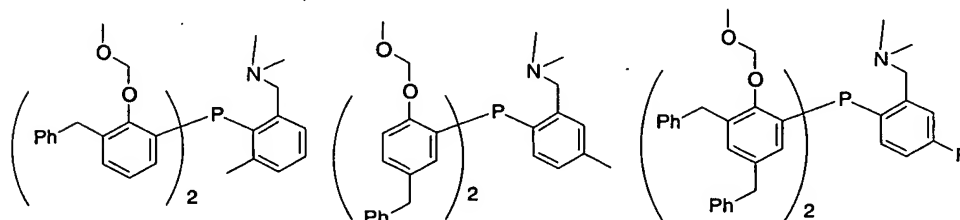
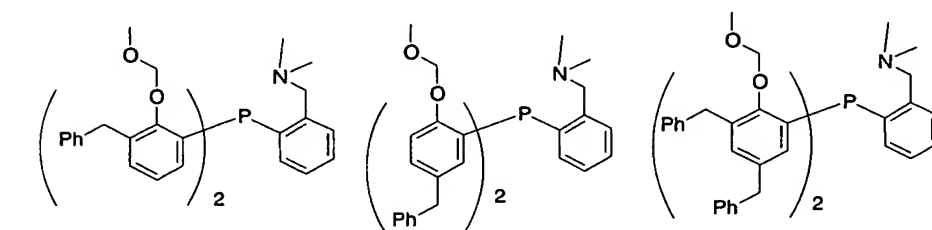
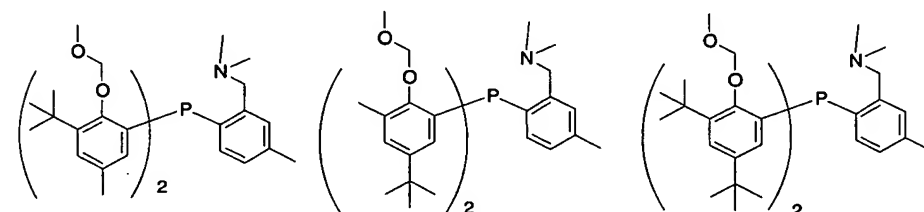
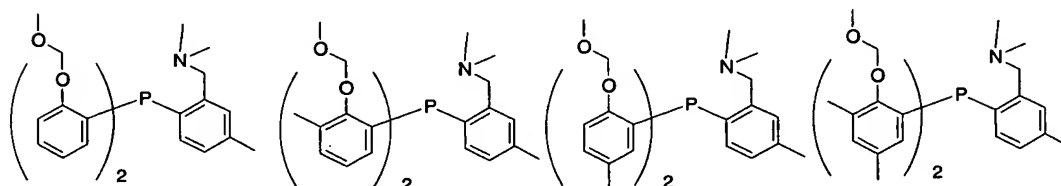


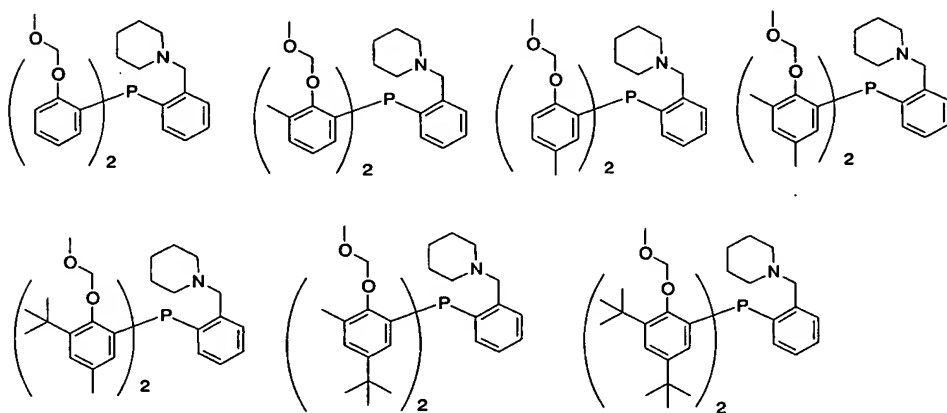
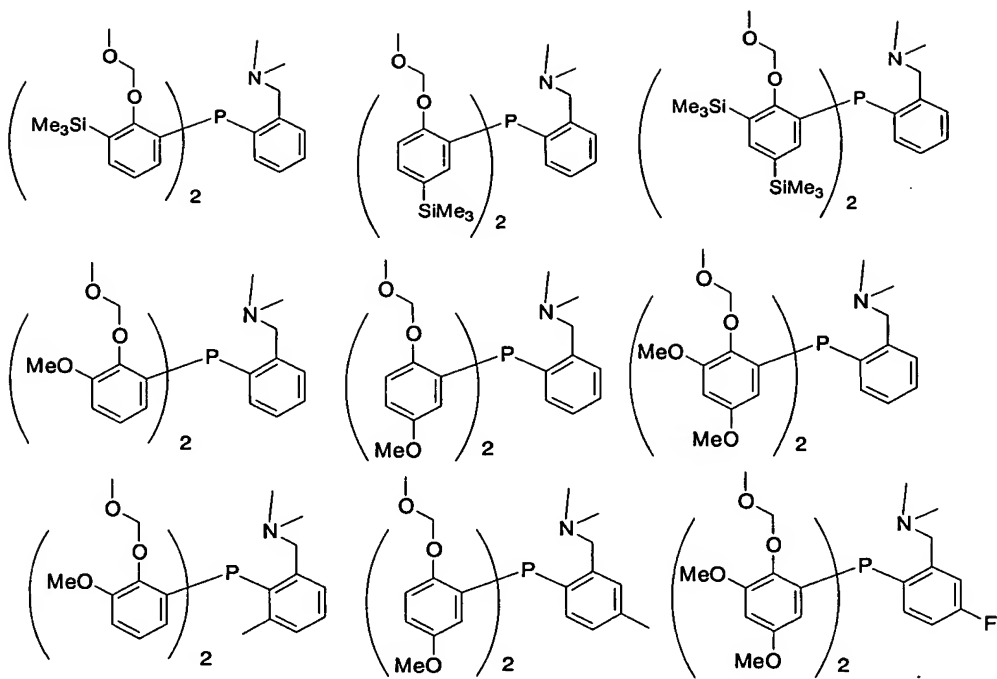


Specific examples of the compound of formula (22B), which corresponds to the compound of formula (1) wherein G^2 is G^{22} include, for example, the following compounds:

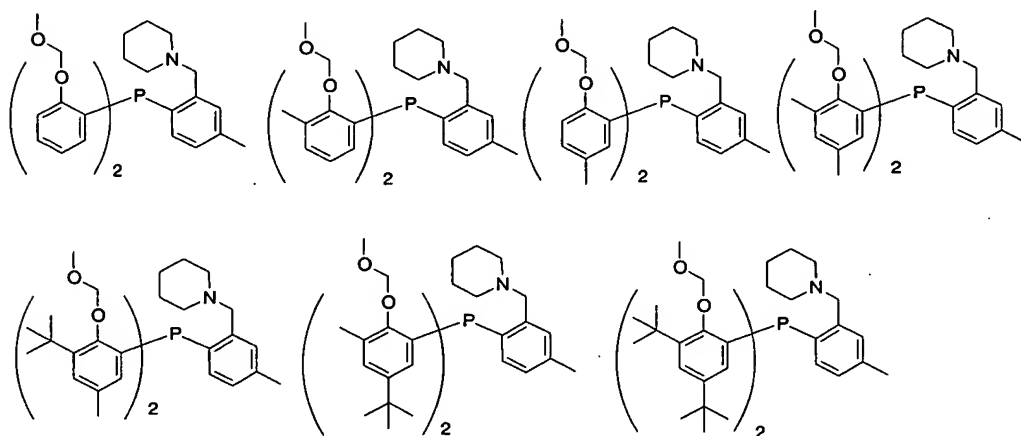


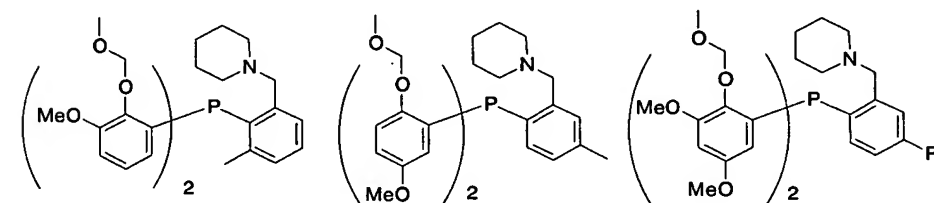
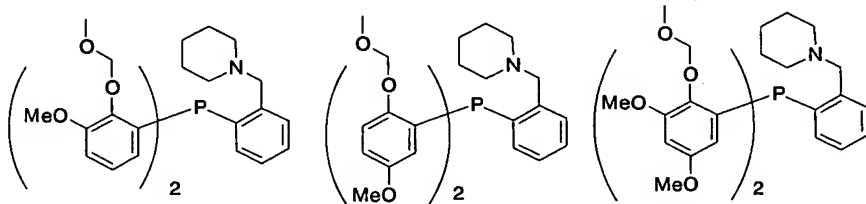
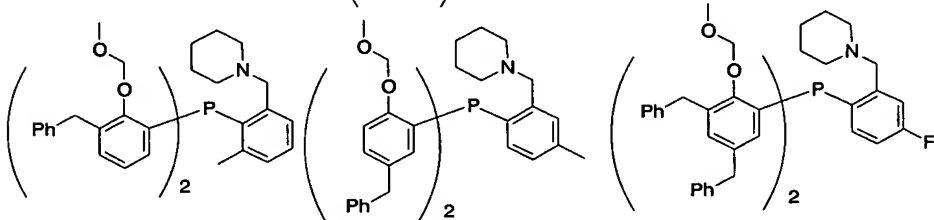
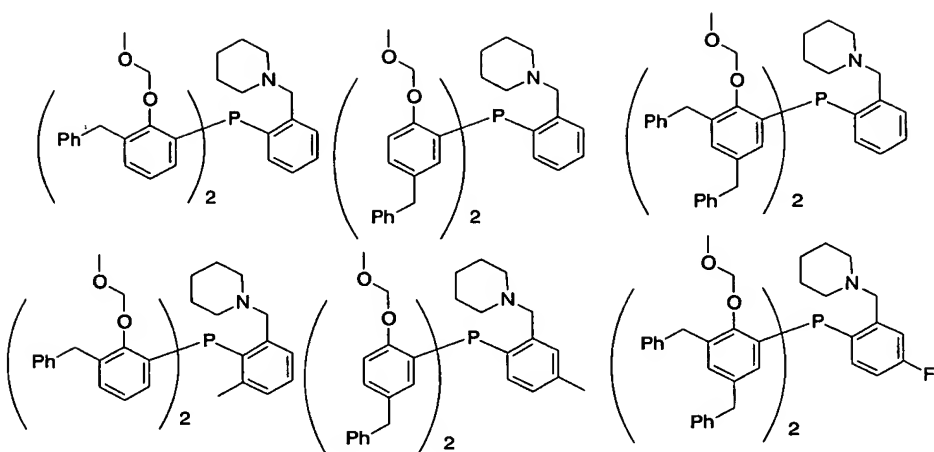
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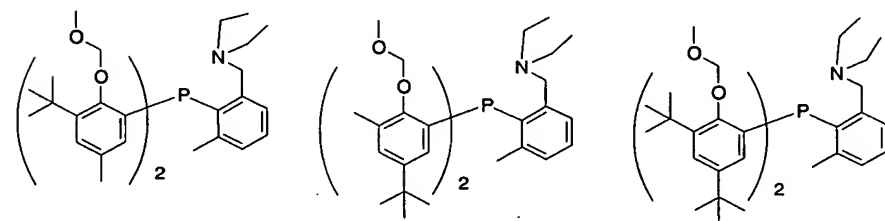
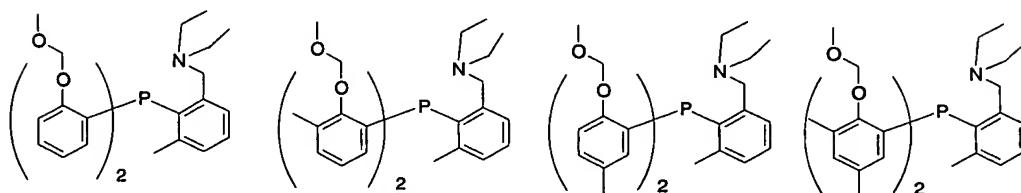


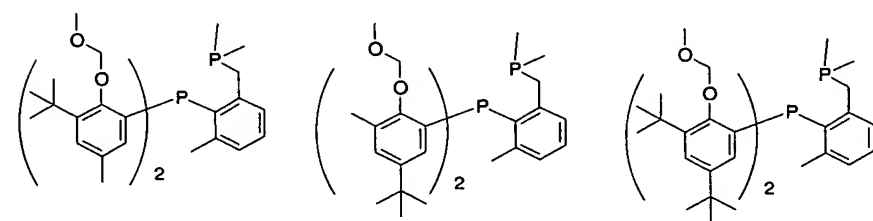
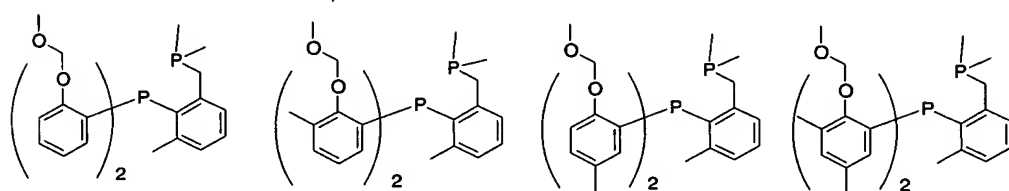
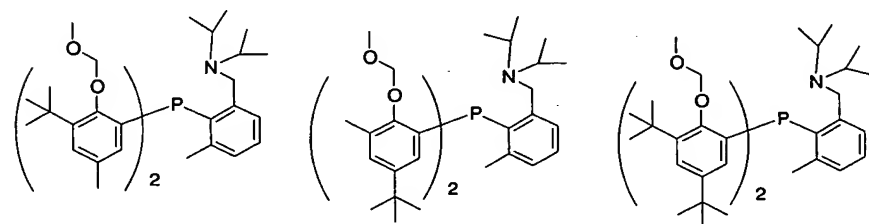
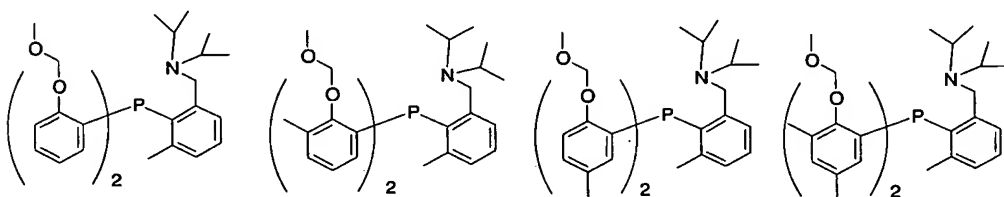
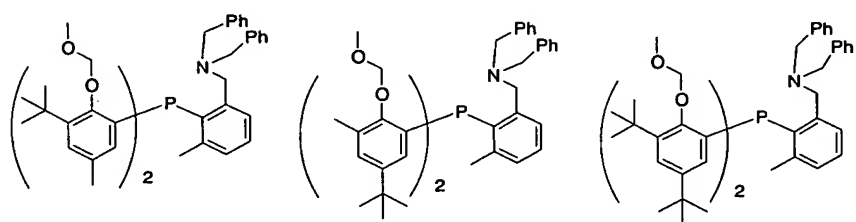
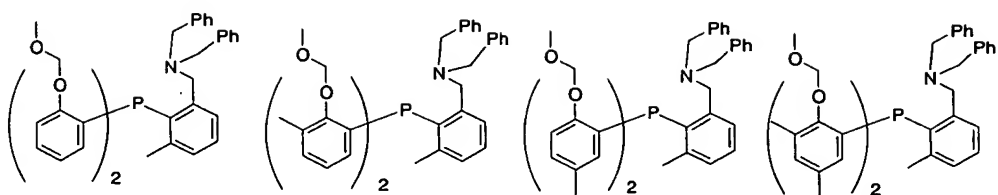
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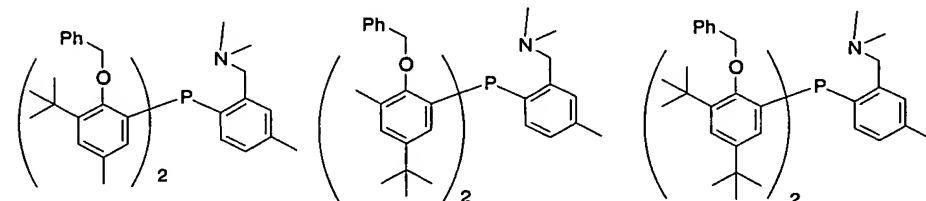
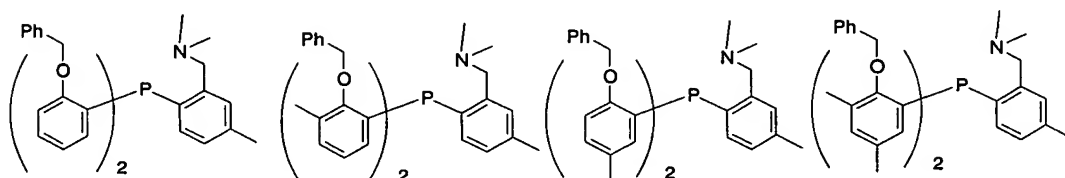
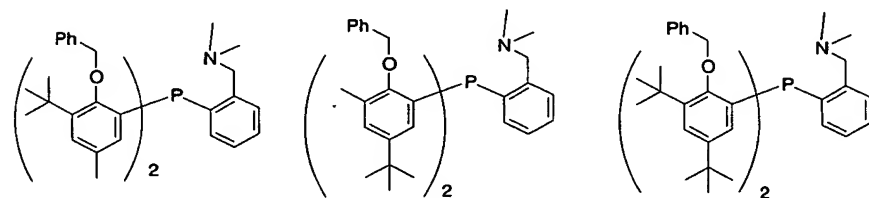
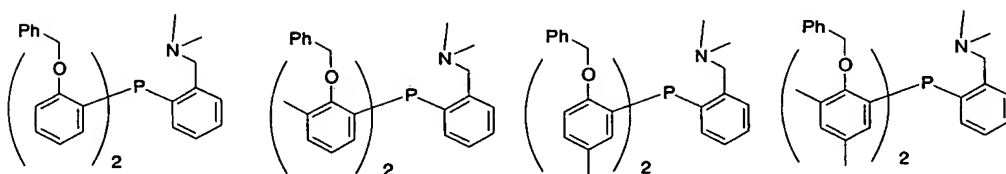
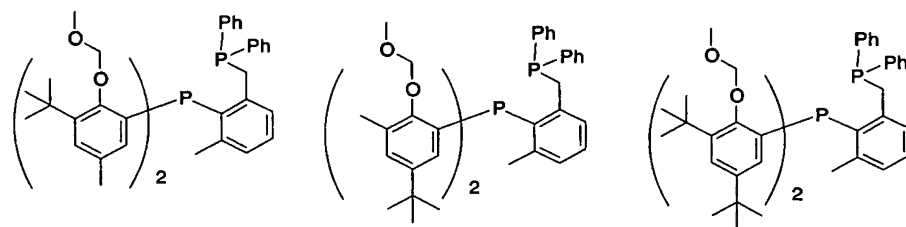
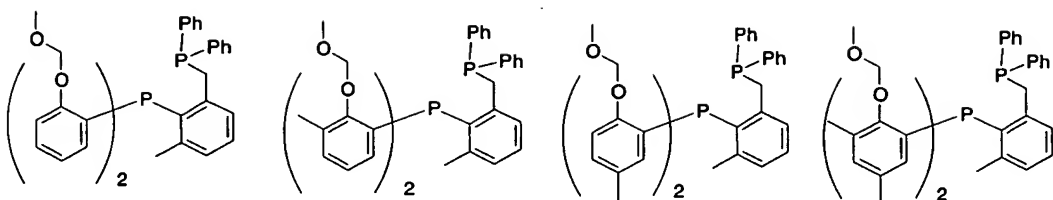


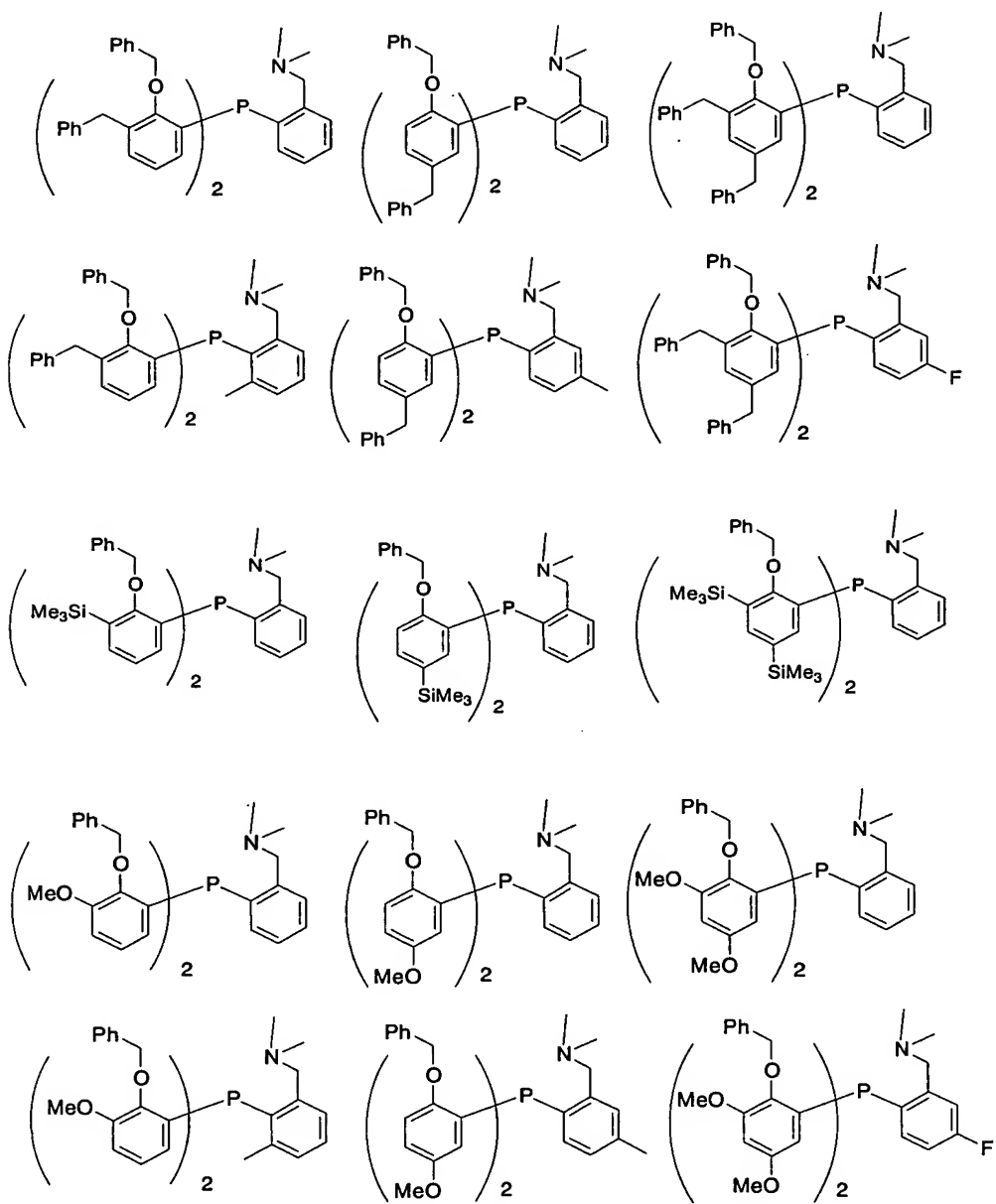


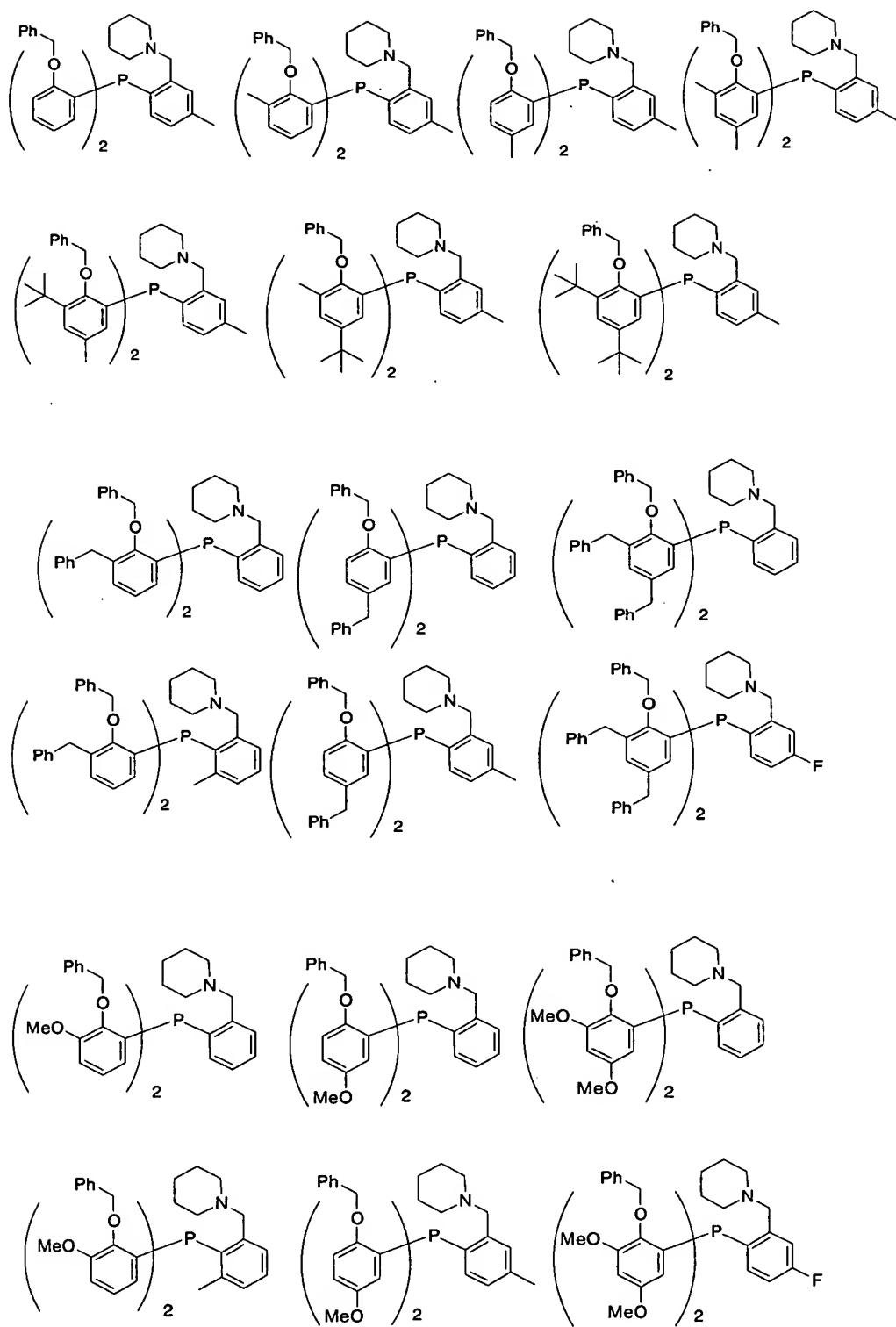
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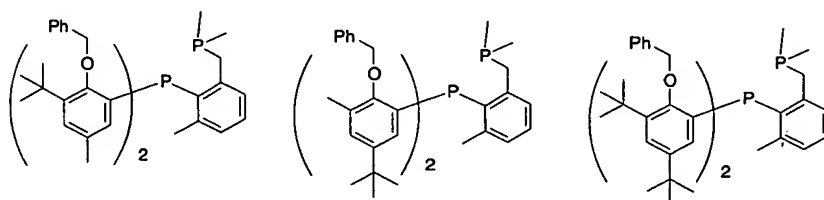
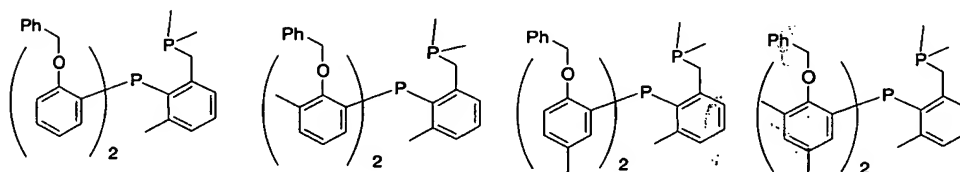
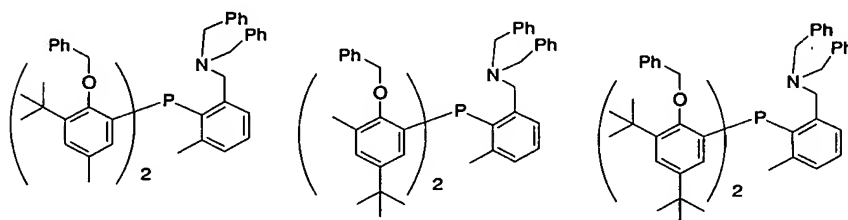
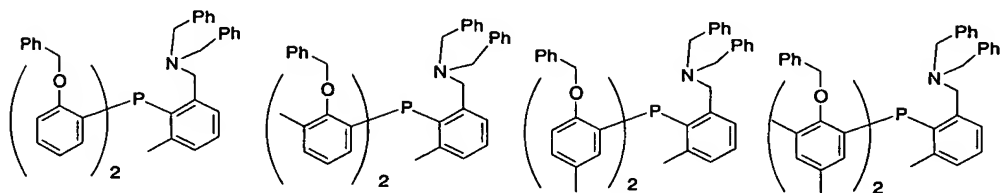
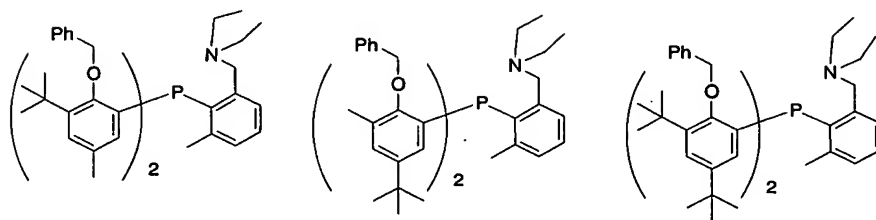
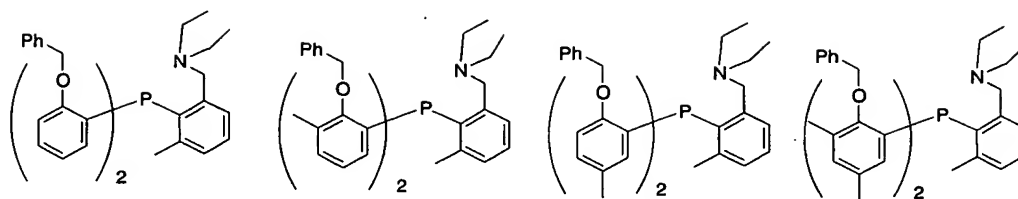


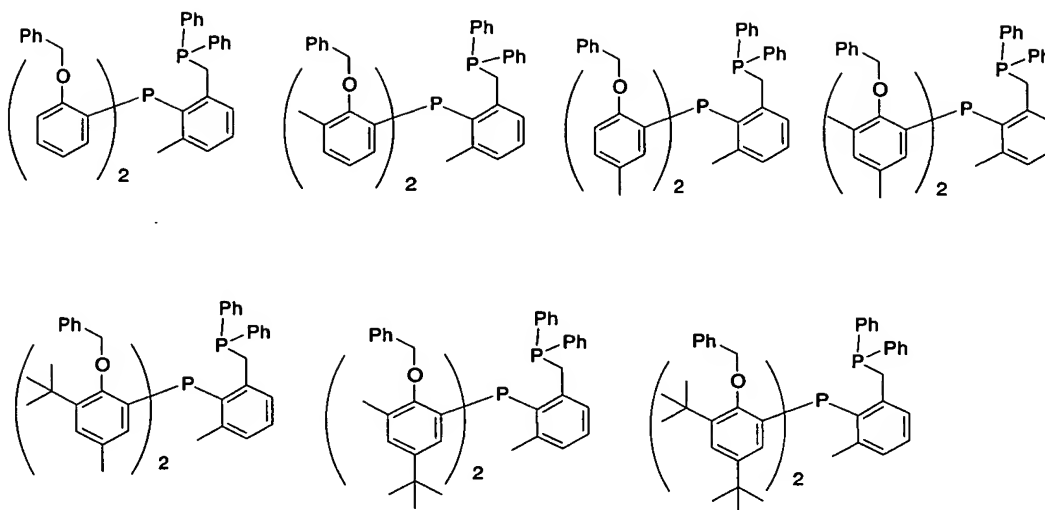












The phosphine compound of formula (22B) can be synthesized by reacting phosphine dihalide of formula (22C) with the metal
 5 aryl compound of formula (22D).

The molar ratio between the phosphine dihalide of formula (22B) and the metal aryl compound of formula (22C) is not particularly restricted, the ratio is preferably in the range of 1:2 to 1:5, more preferably in the range of 1:2 to 1:2.5.

10 Specific examples of the halogen atom represented by X^2 include fluorine, chlorine, bromine and iodine atoms, and the chlorine atom is preferable.

Specific examples of the alkali metal and the alkaline earth metal represented by D in formula (22D) include lithium,
 15 sodium, potassium, magnesium and calcium atoms, and lithium and magnesium atoms are preferable.

While the reaction is usually carried out in a solvent inert to the reaction, examples of the solvent include aromatic hydrocarbon solvents such as benzene, toluene or the like,

aliphatic hydrocarbon solvents such as hexane, heptane or the like, and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture of at least two of them, and the ratio of use thereof is usually in the range of 1 to 200 parts by weight, preferably in the range of 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (22D).

The reaction can be usually performed by adding phosphine dihalide of formula (22C) to the metal aryl compound of formula (22D). The reaction temperature is usually in the range of from -100°C or more to the boiling temperature of the solvent or less, preferably in the range of -80°C to 100°C.

The phosphine compound of formula (22B) can be obtained by removing insolubles by a conventional method such as filtration, and by removing the solvent by evaporation. The product is purified by silica gel column chromatography, if necessary.

Phosphine dihalide of formula (22C) is produced by a reaction between trihalogenated phosphorous of formula:

$P(X^2)_3$ (X^2 represents a halogen atom) and a metal aryl compound.

For example, phosphine dihalide may be produced by a reaction between the metal aryl compound of formula (22E) and phosphorous trihalide.

The molar ratio between the metal aryl compound of formula (22E) and phosphorous trihalide that may be used in the reaction is not particularly restricted, and it is preferably in the range of 1:1 to 1:5, more preferably in the range of 1:1 to 1:2.5.

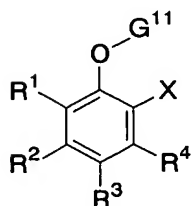
The reaction is usually performed in a solvent inert to

the reaction. Examples of the solvent include aromatic hydrocarbon solvents such as benzene, toluene or the like, aliphatic hydrocarbon solvents such as hexane, heptane or the like, and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture of at least two of them, and the ratio thereof that may be used is usually in the range of 1 to 200 parts by weight, preferably in the range of 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (22E).

The reaction can be usually performed by adding phosphorous trihalide to the metal aryl compound of formula (22E). The reaction temperature is usually in the range from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C.

The phosphine dihalide of formula 22C can be obtained from the reaction mixture obtained by a conventional method, for example by removing insoluble products by filtration followed by removing the solvent by evaporation. The product may be purified by distillation, if necessary.

The metal aryl compound of formula (22D) can be produced by reacting an organic compound of formula (22F):



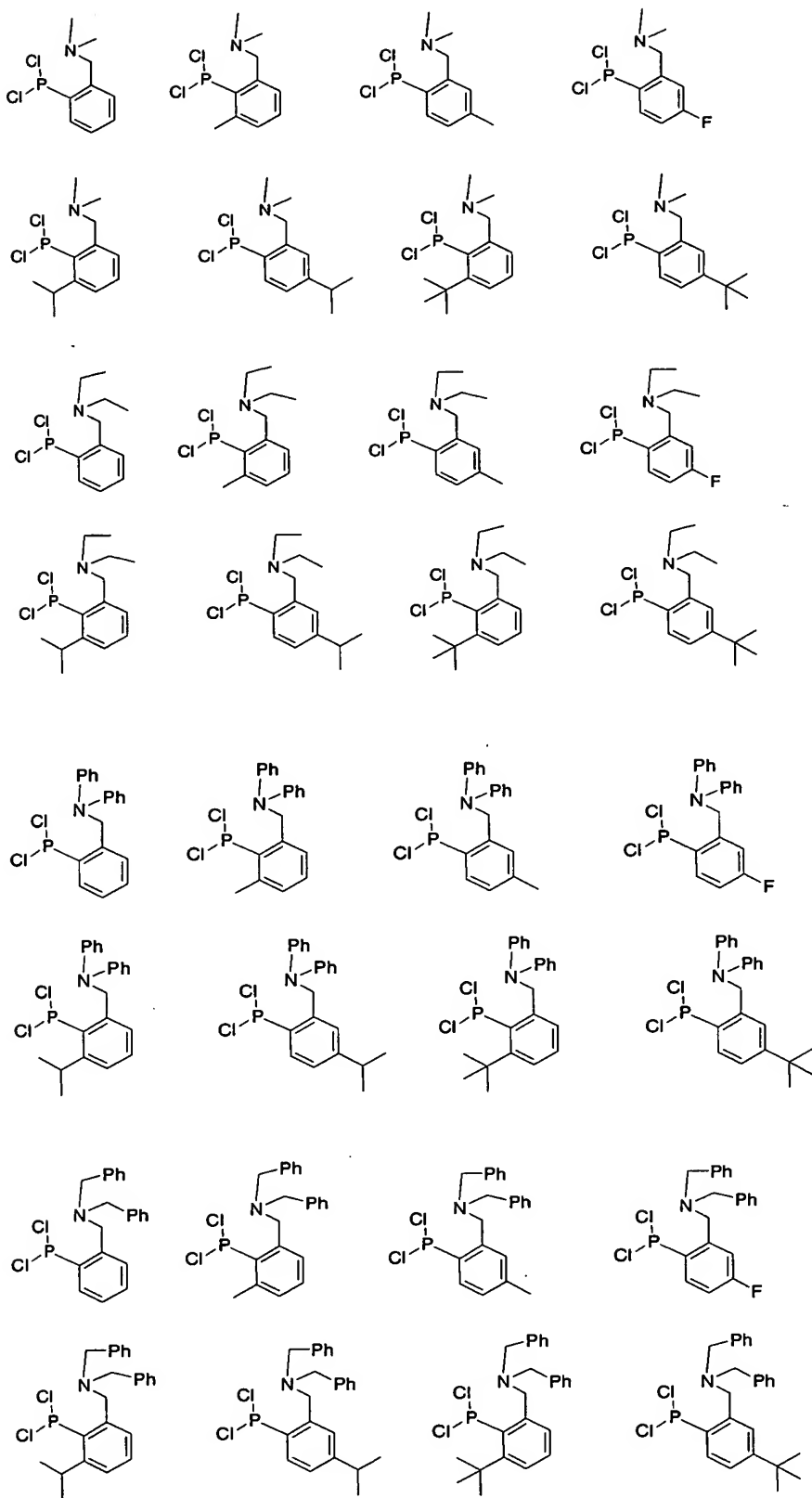
(22F)

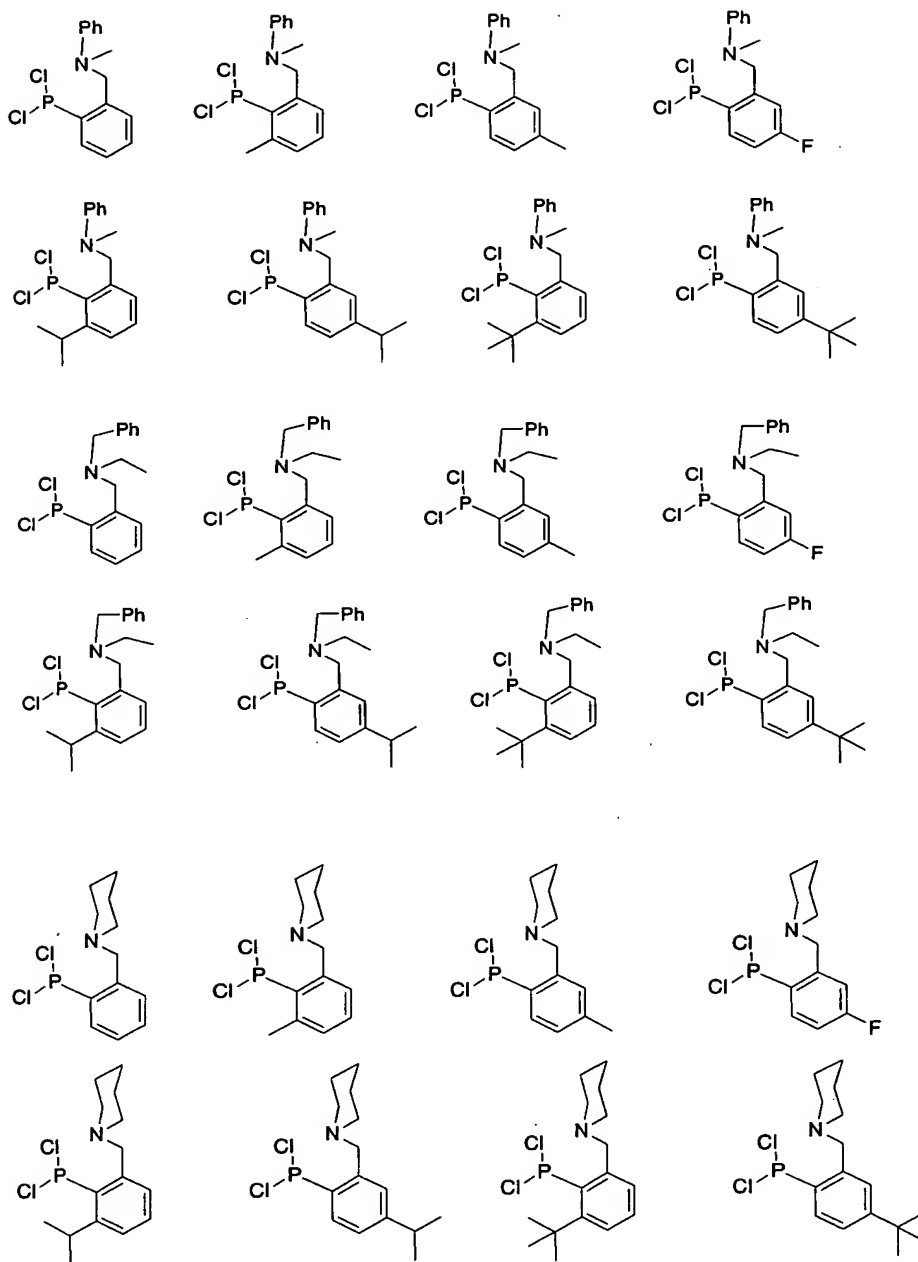
wherein R¹, R², R³, R⁴ and G¹¹ each denotes the same meaning as described above, and X represents a hydrogen atom or a halogen atom, with a lithiating agent when X is hydrogen, or with the

lithiating agent or magnesium metal, when X is a halogen atom. The molar ratio between the organic compound of formula (22F) and the lithiating agent or magnesium metal that may be used in the reaction is not particularly restricted, and it is
5 preferably in the range of 1:1 to 1:5, more preferably in the range of 1:1 to 1:2.5. Examples of the lithiating agent include methyl lithium, n-butyl lithium, s-butyl lithium, t-butyl lithium and phenyl lithium, and preferred is n-butyl lithium.

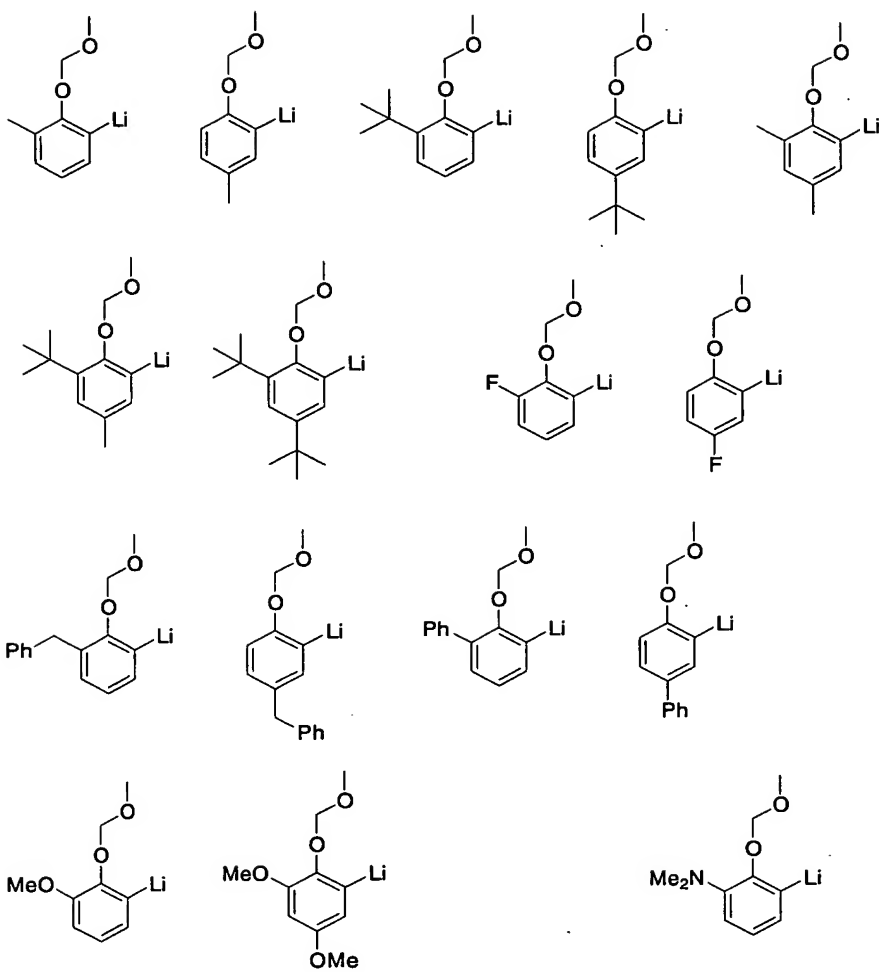
The reaction is usually performed in a solvent inert to
10 the reaction. Examples of the solvent include aromatic hydrocarbon solvents such as benzene, toluene or the like, aliphatic hydrocarbon solvents such as hexane, heptane or the like, and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture
15 of at least two of them, and the ratio thereof is usually in the range of 1 to 200 parts by weight, preferably in the range of 3 to 50 parts by weight, per part by weight of the organic compound of formula (22F). The reaction can be usually performed by adding the lithiating agent to organic compound of formula
20 (22F). The reaction temperature is usually in the range from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C.

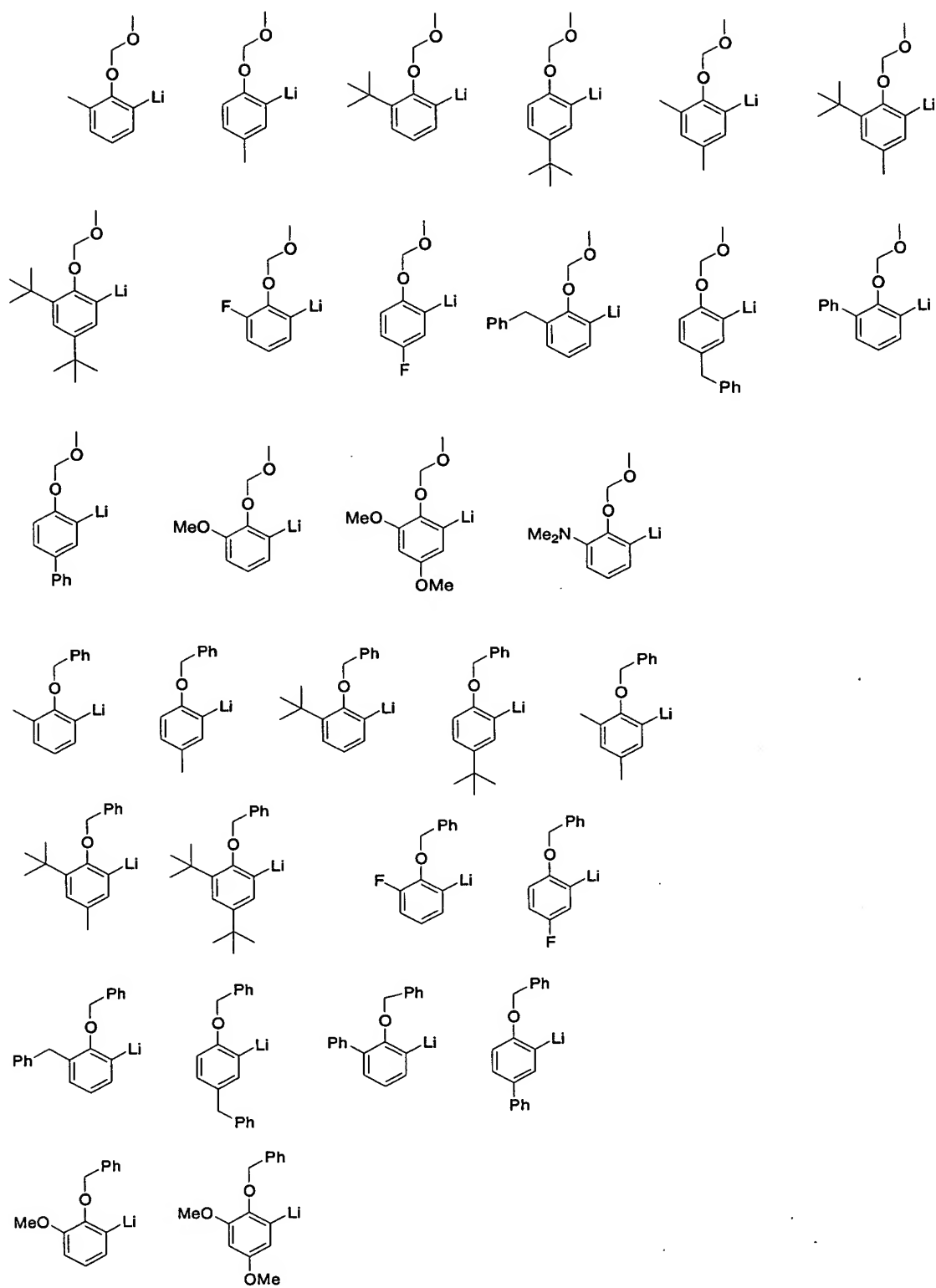
Specific examples of phosphine dihalide of formula (22C) include, for example, the following compounds and those that
25 have bromine or iodine atom in place of the chlorine atom in the illustrated compounds:

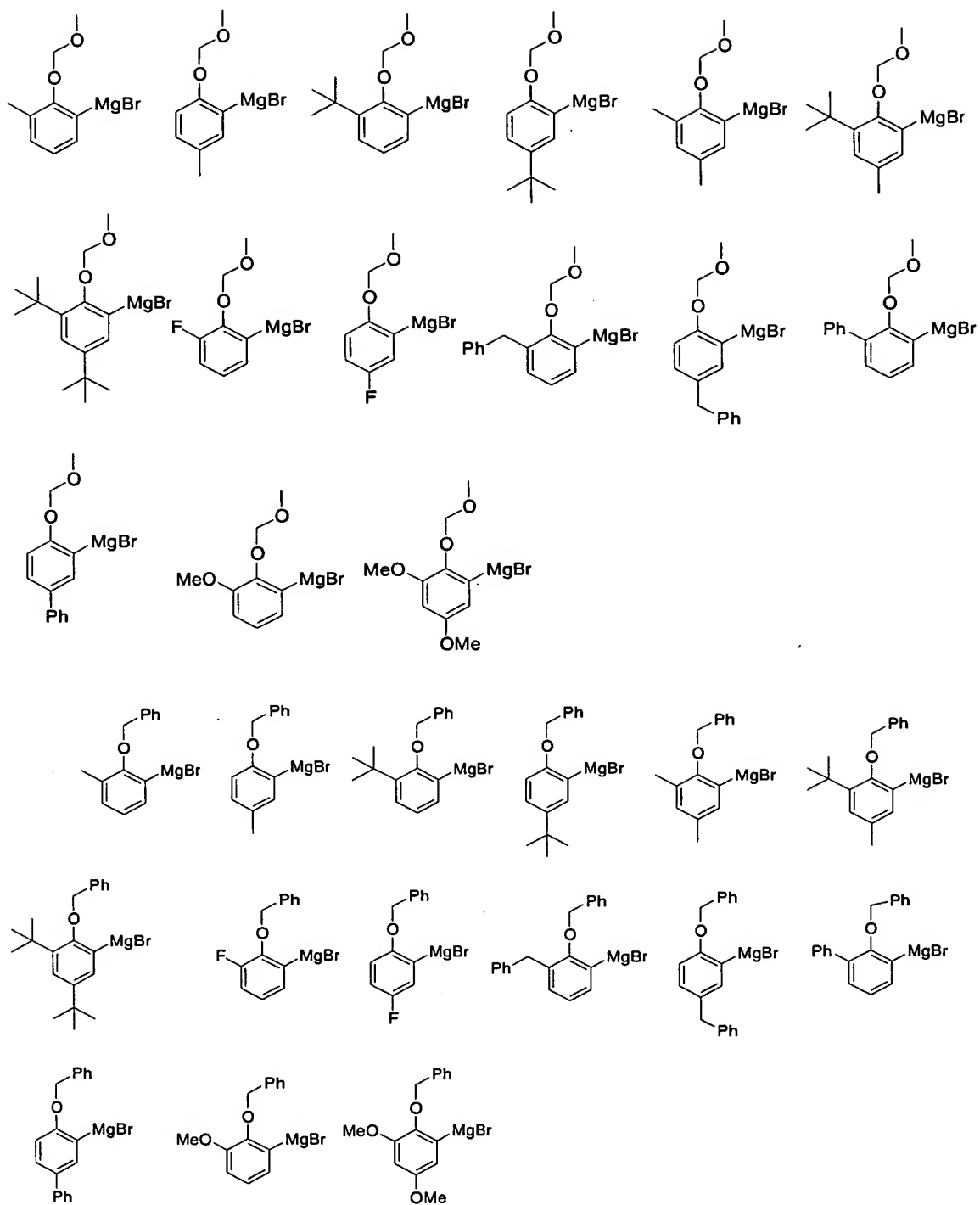




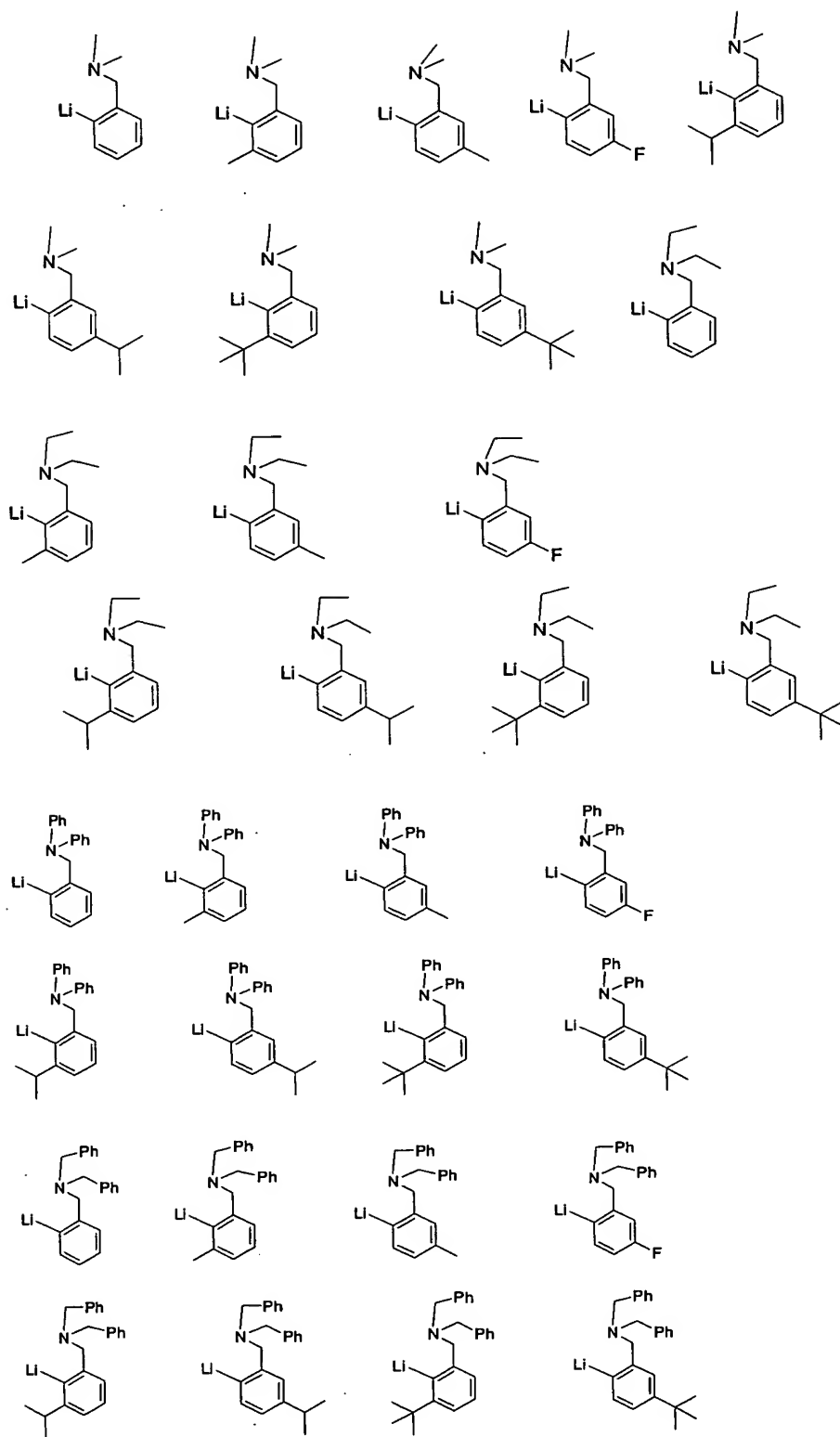
Specific examples of the metal aryl compound of formula (22D) include, for example, the following compounds:

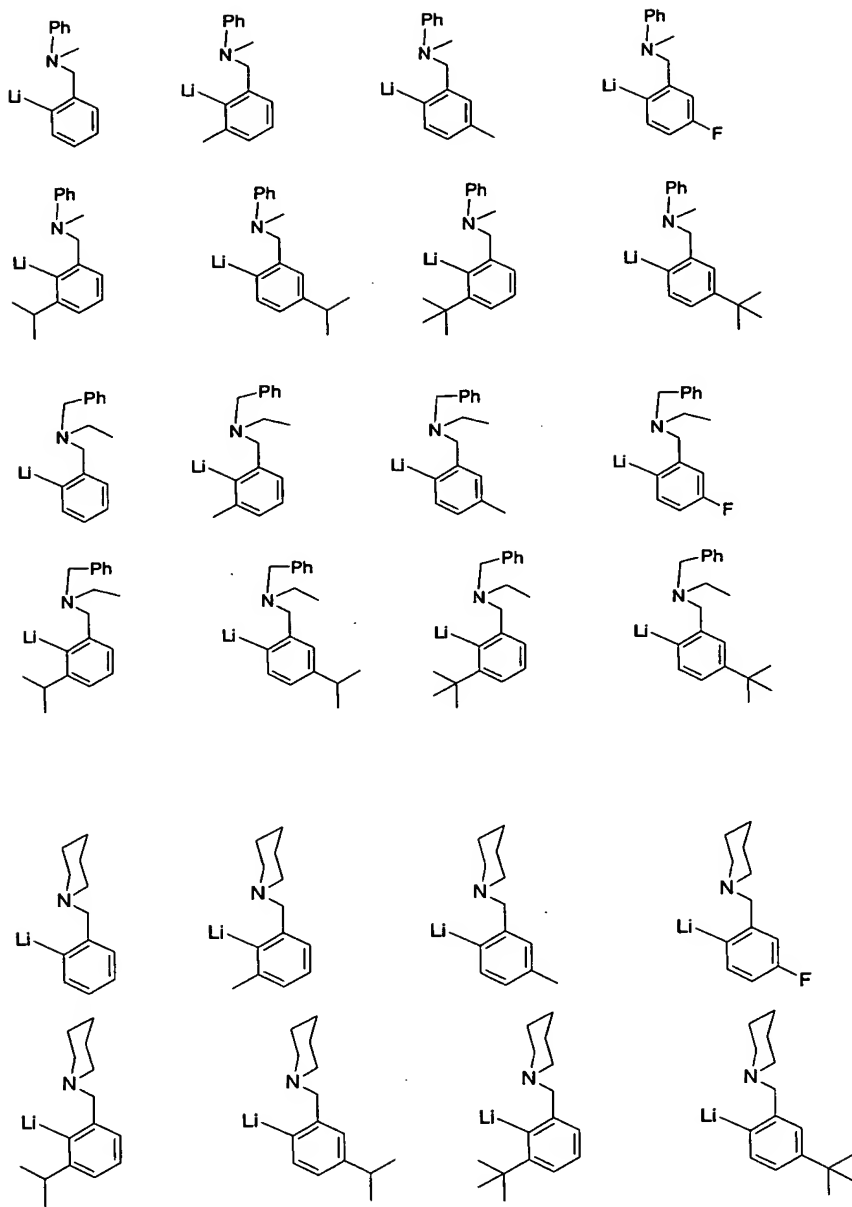


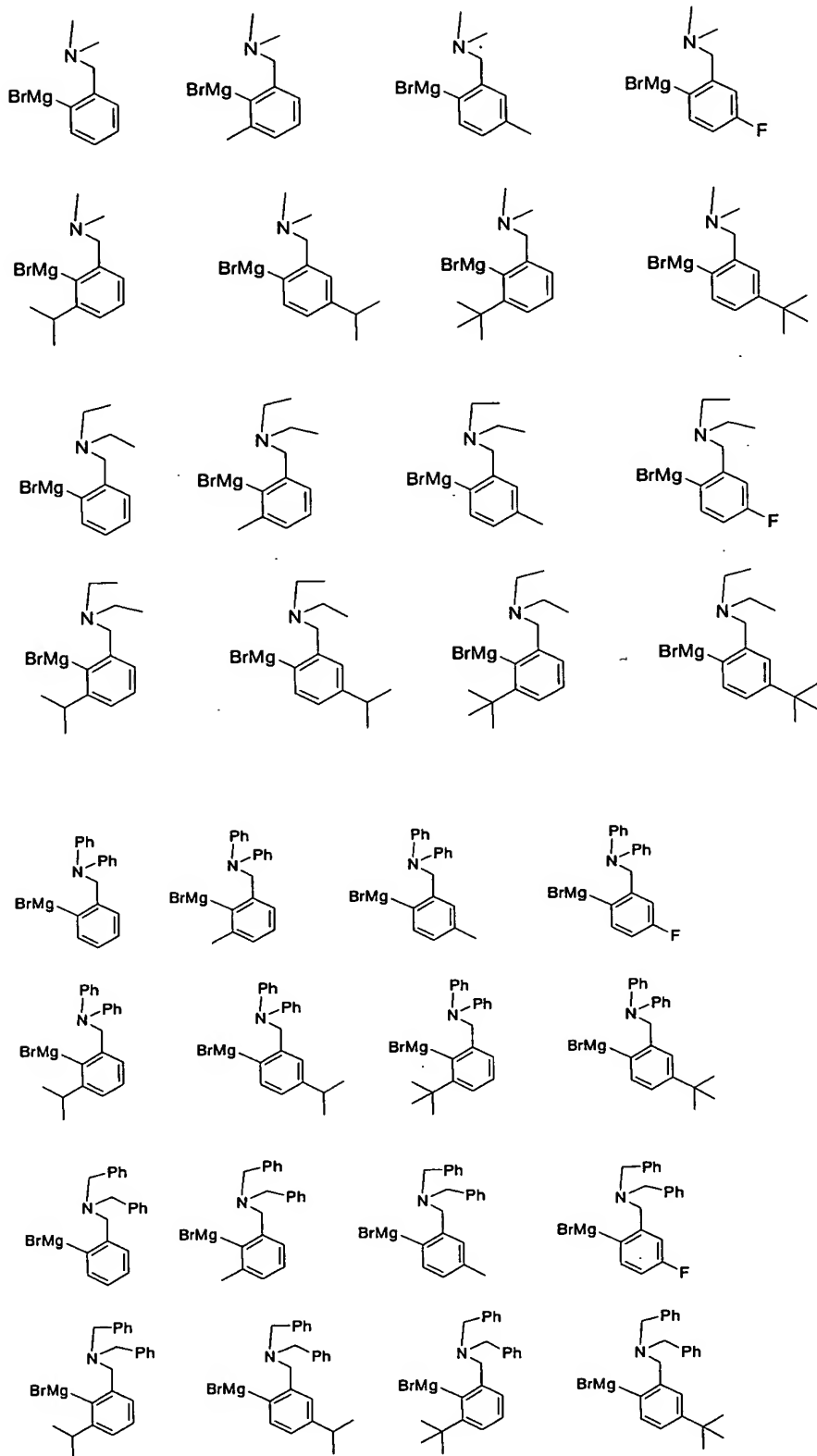


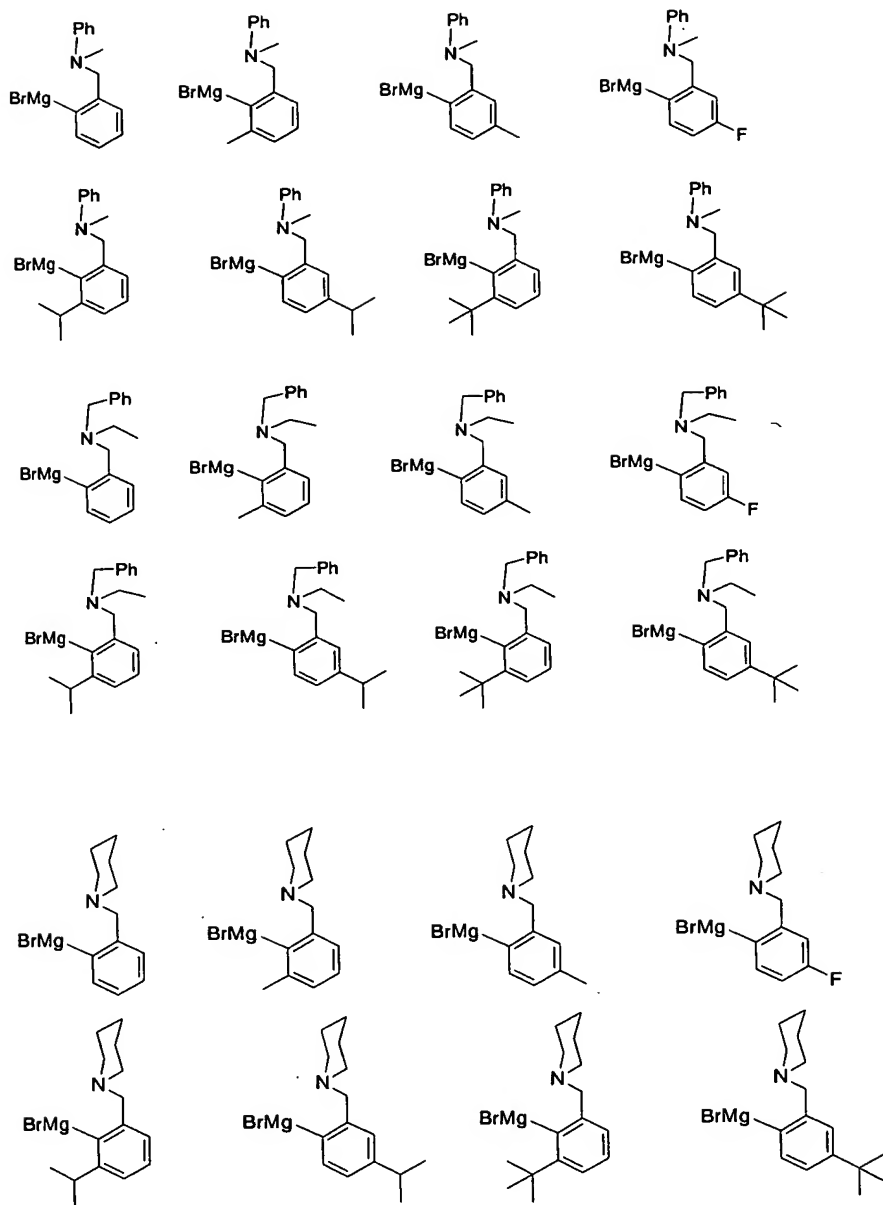


5 Specific examples of the metal aryl compound of formula (22E) include, for example, the following compounds:

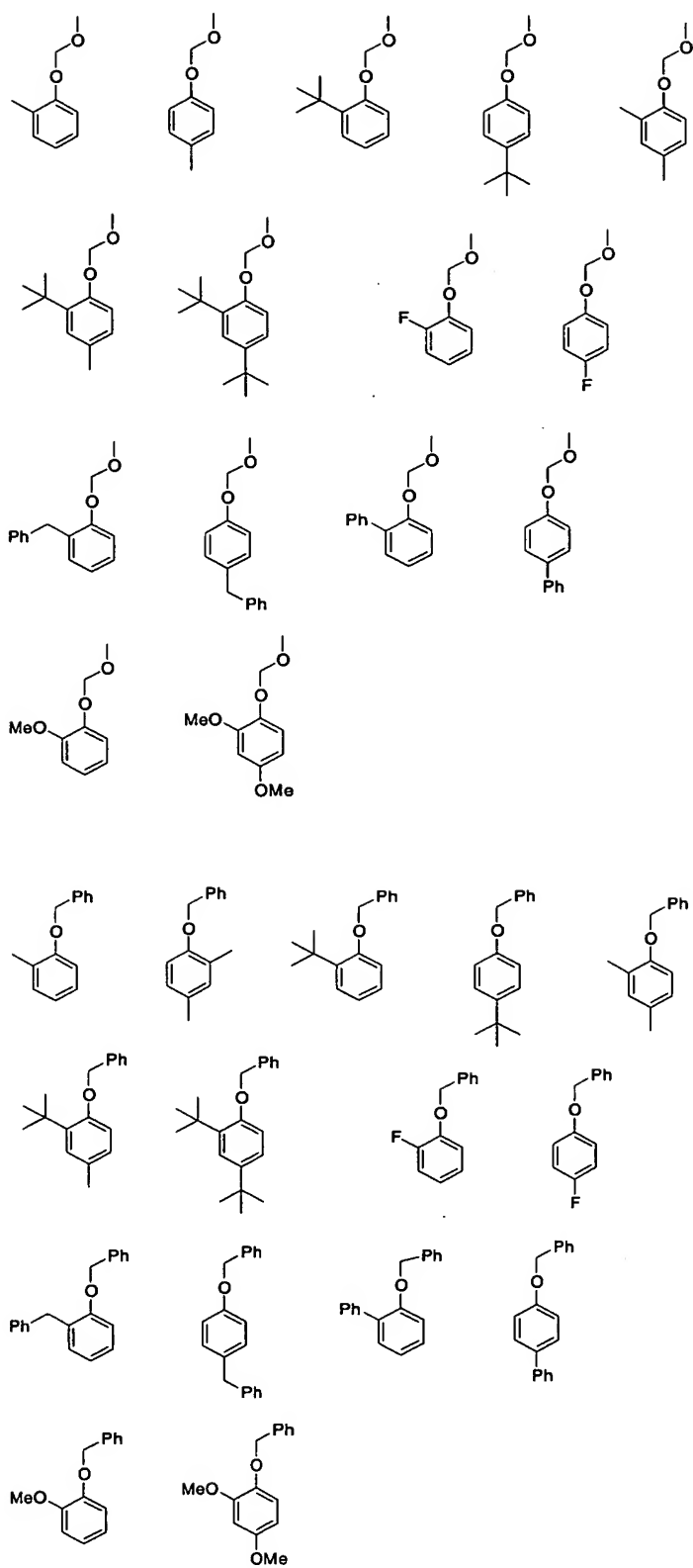


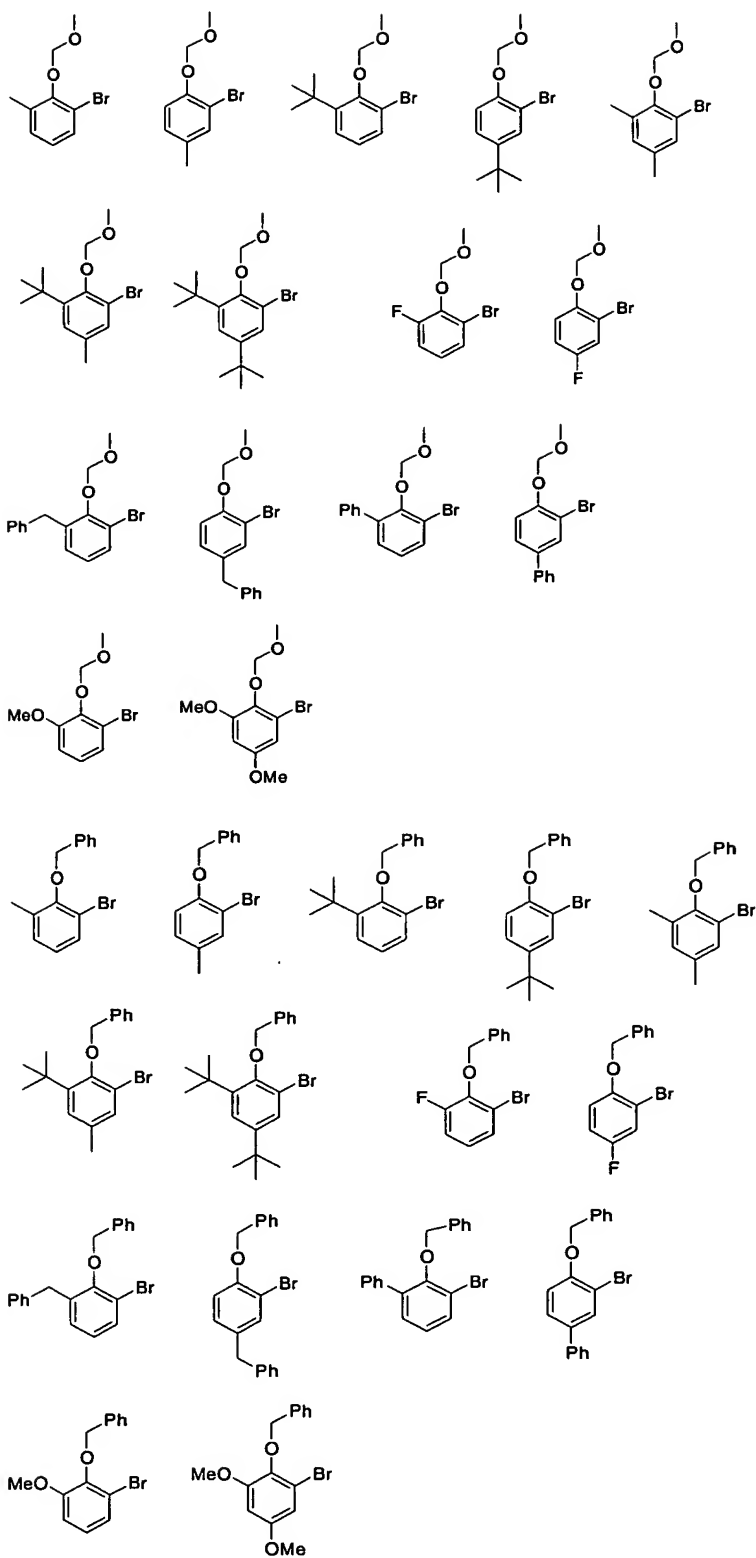






Specific examples of the starting material for the compound
 5 of formula (22E) include, for example, the following compounds:

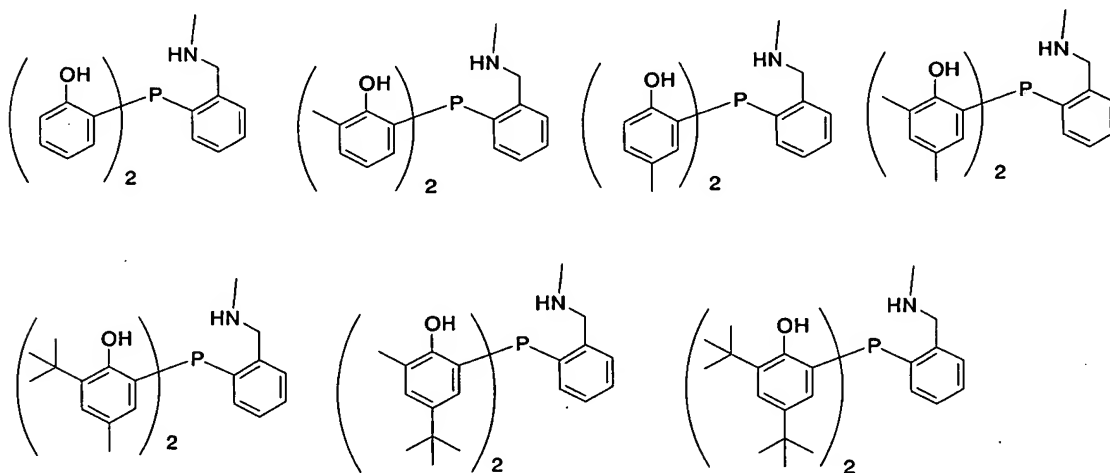


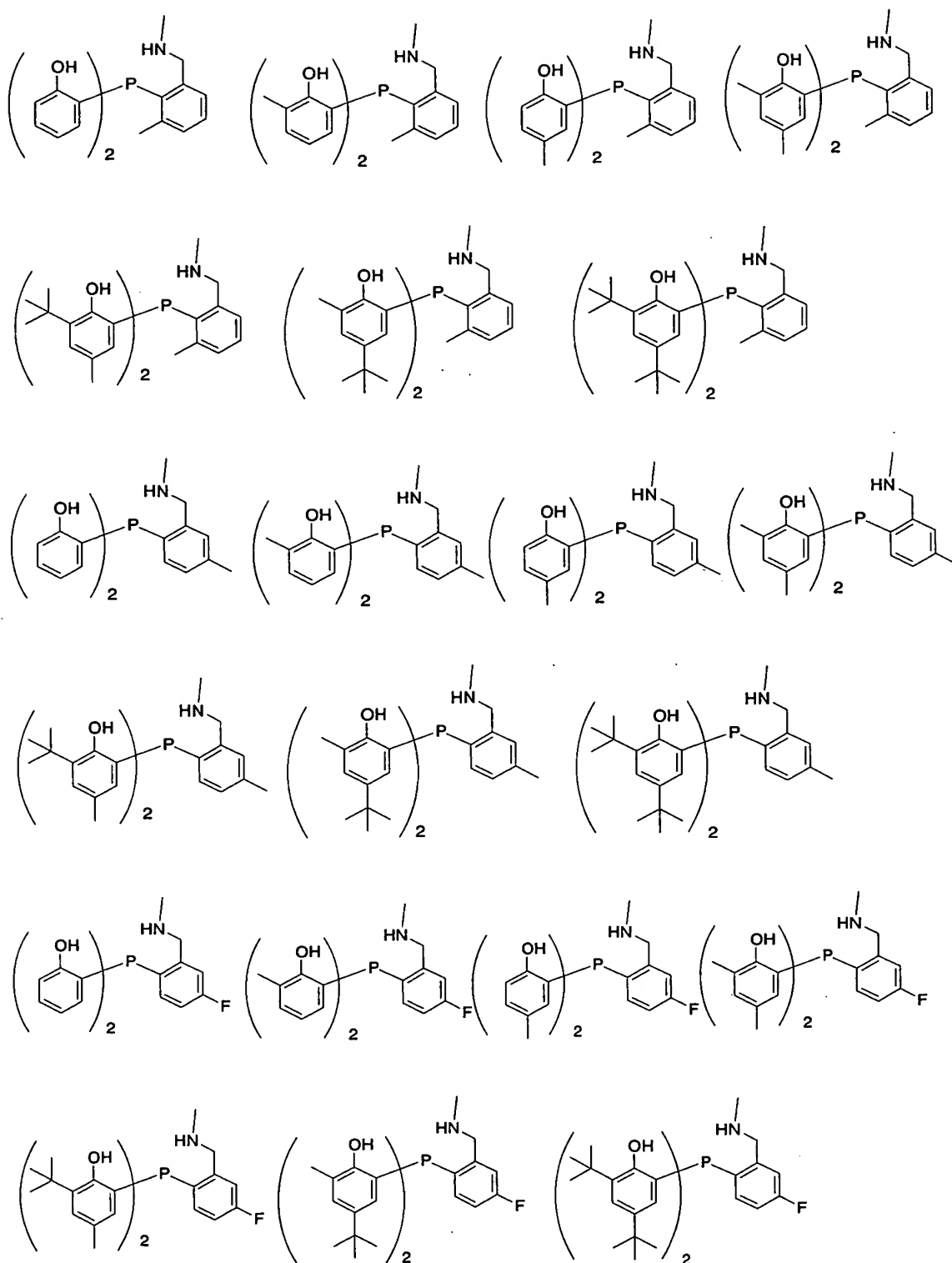


The compound in which A^1 is P in the compound of formula (22A) can be synthesized by deprotection the compound of formula (23B) wherein A^1 is P as in the compound in which A^1 is N. The compound of formula (23B) wherein A^1 is P can be synthesized by using the compound of formula (22E) wherein A^1 is P as a starting material. An excess amount of an alkali metal reagent may be used in the coupling reaction by suitably adjusting, if necessary.

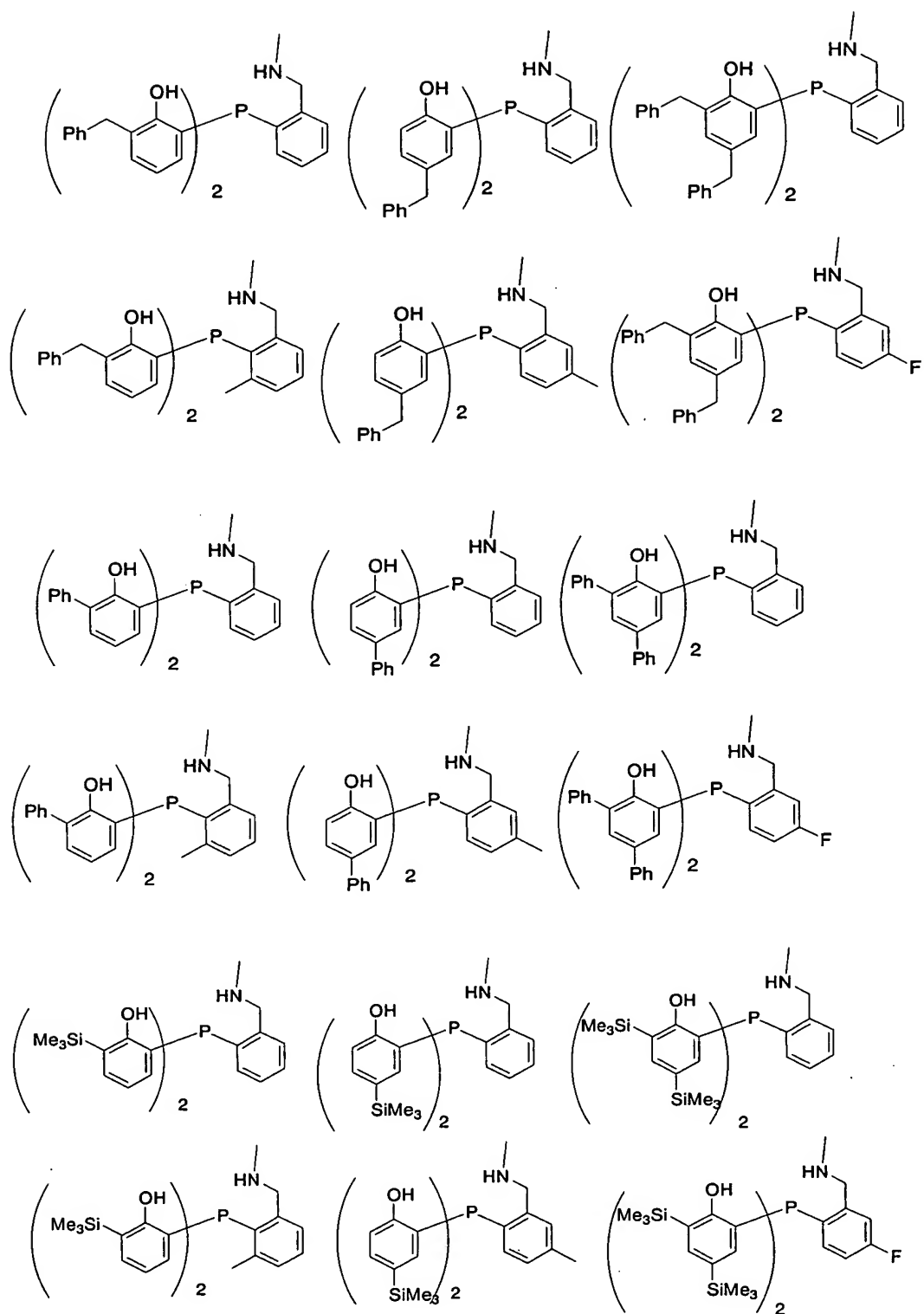
The halogenated aryl compound as a precursor of the compound in which A^1 is P in formula 22E may be synthesized by a known method described (for example, Zeitschrift fuer Anorganische Chemie, vol. 494, p55, 1892), for example by activating 1-(α -bromomethyl)-2-bromobenzene with an alkali metal, followed by reacting a chlorophosphine compound.

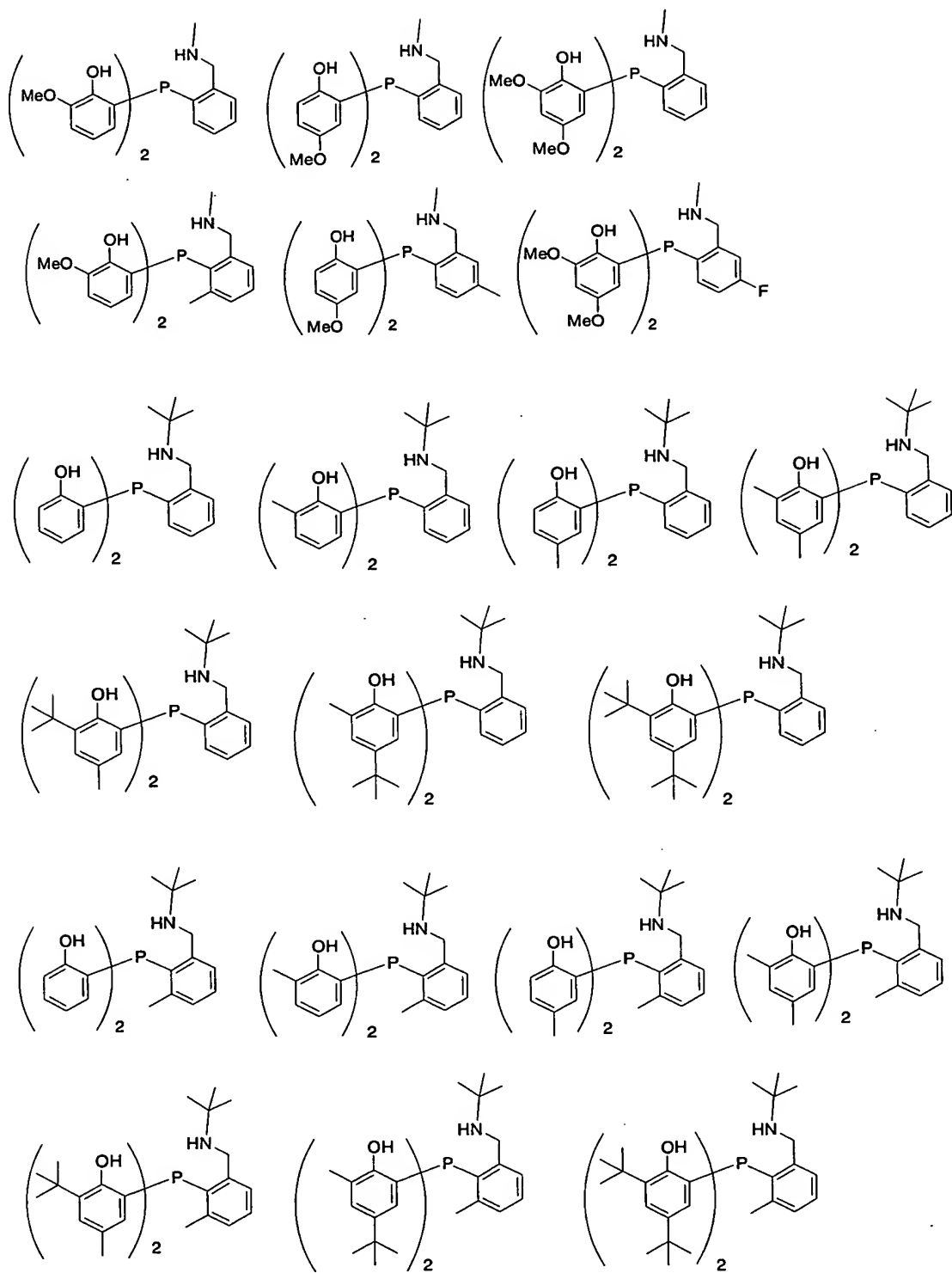
Examples of the compound of formula 23A, which corresponds to the compounds of formula (1) wherein G^1 has the structure of G^{23} will be shown below:

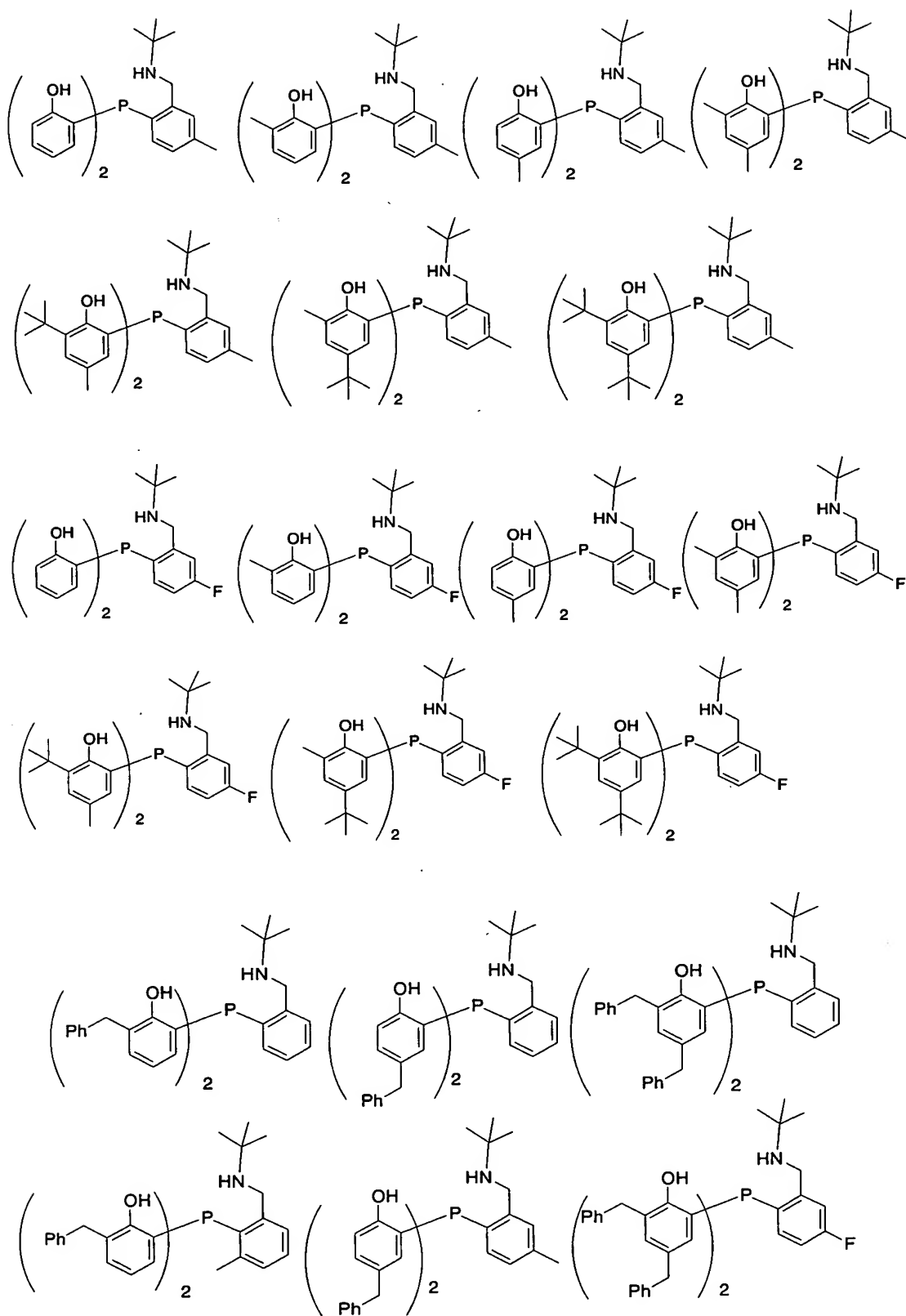


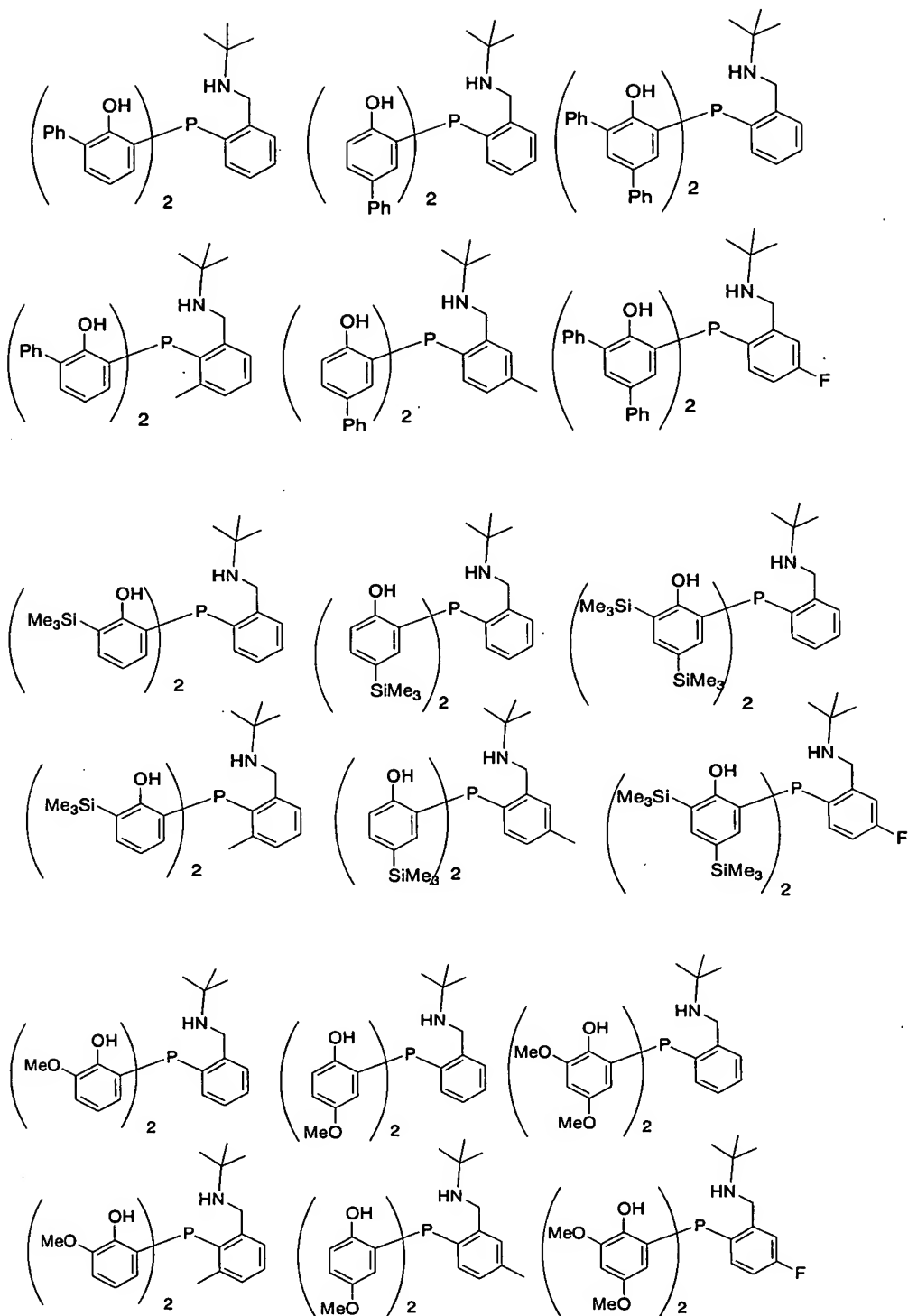


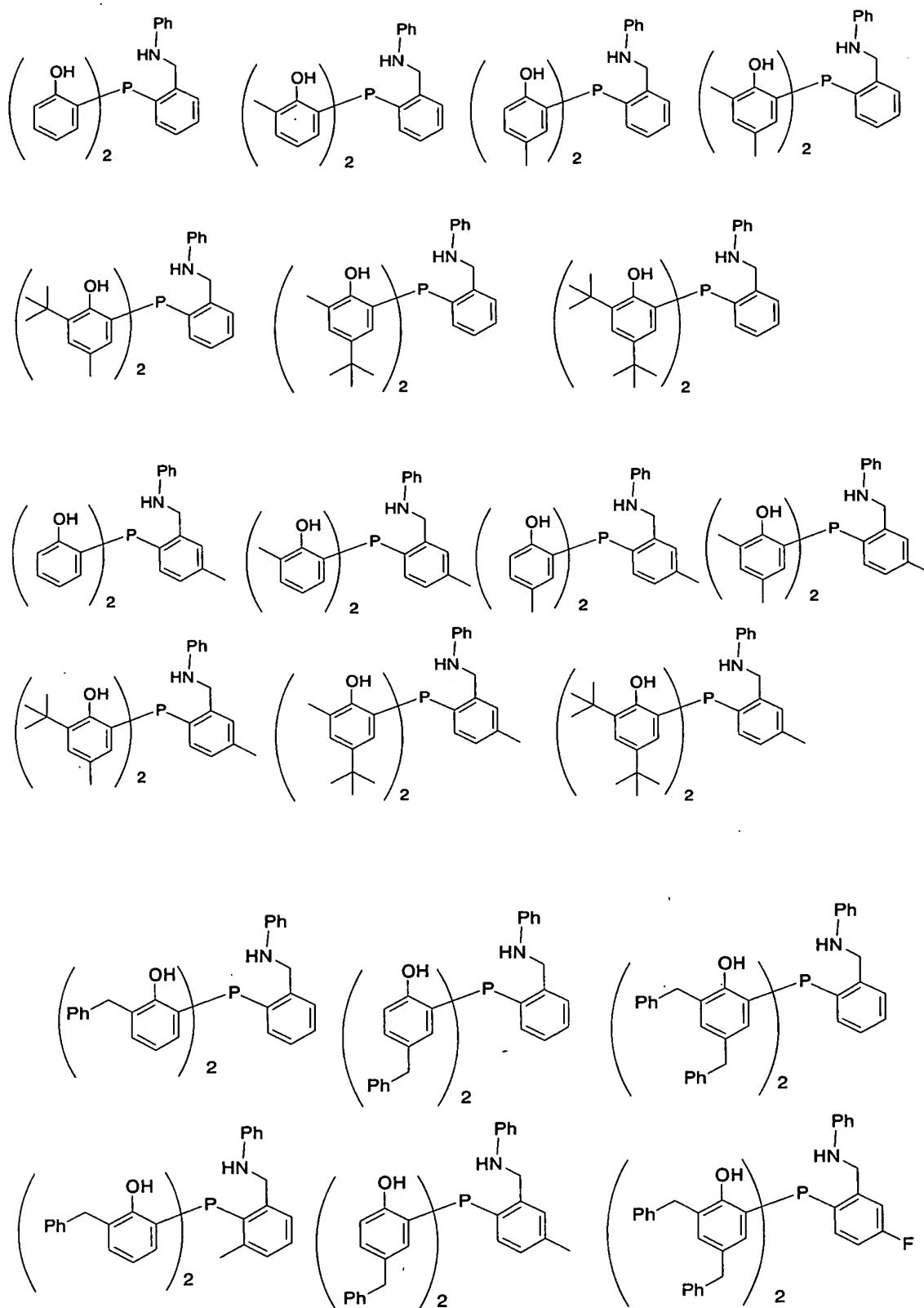
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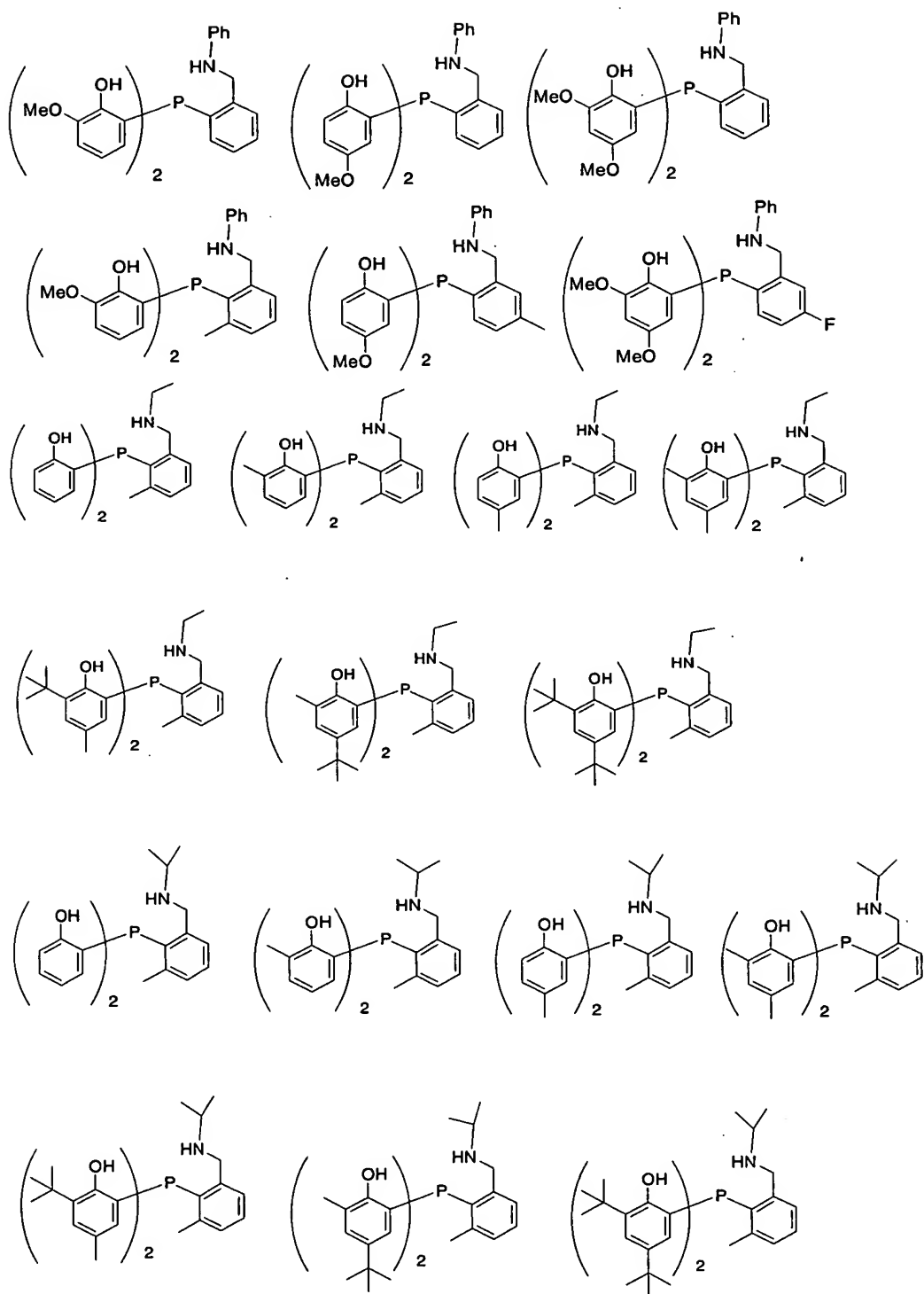


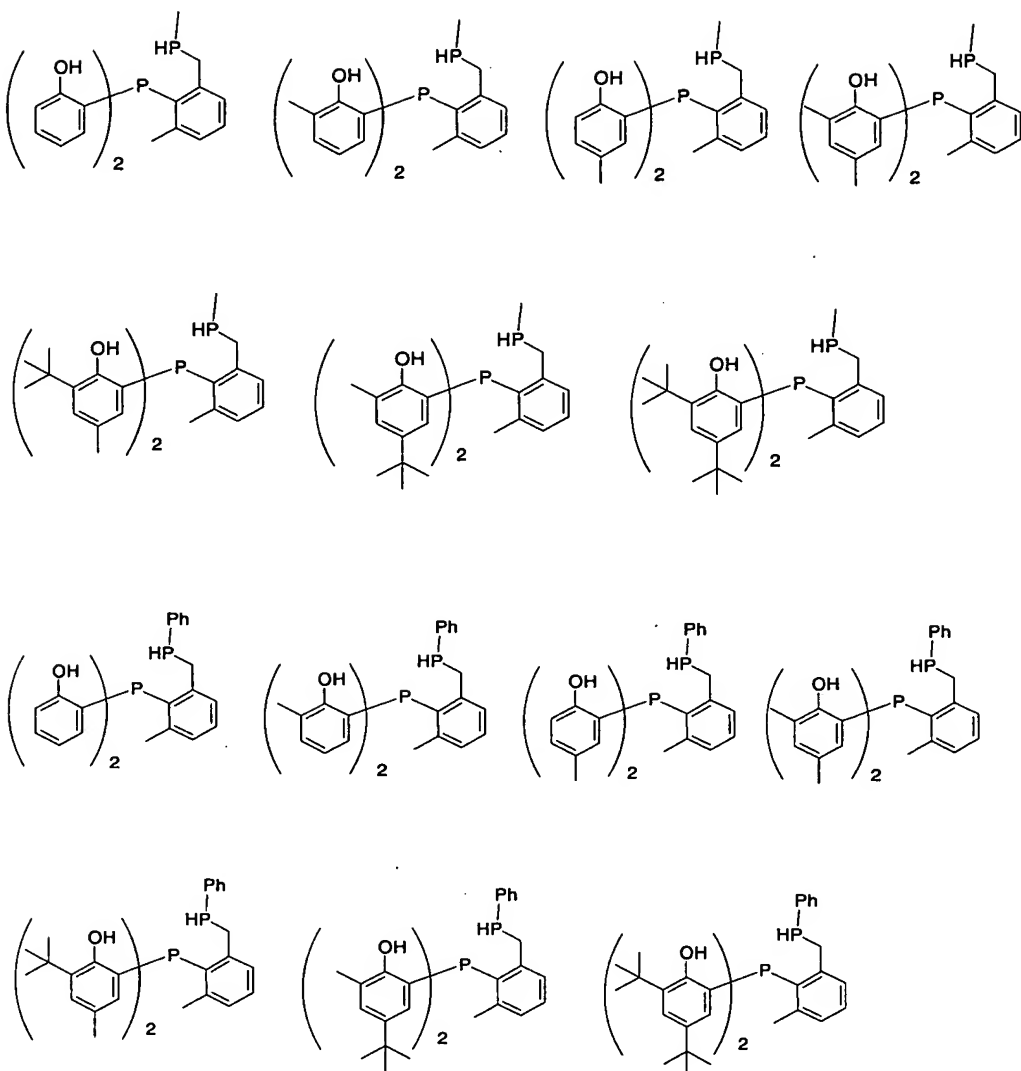




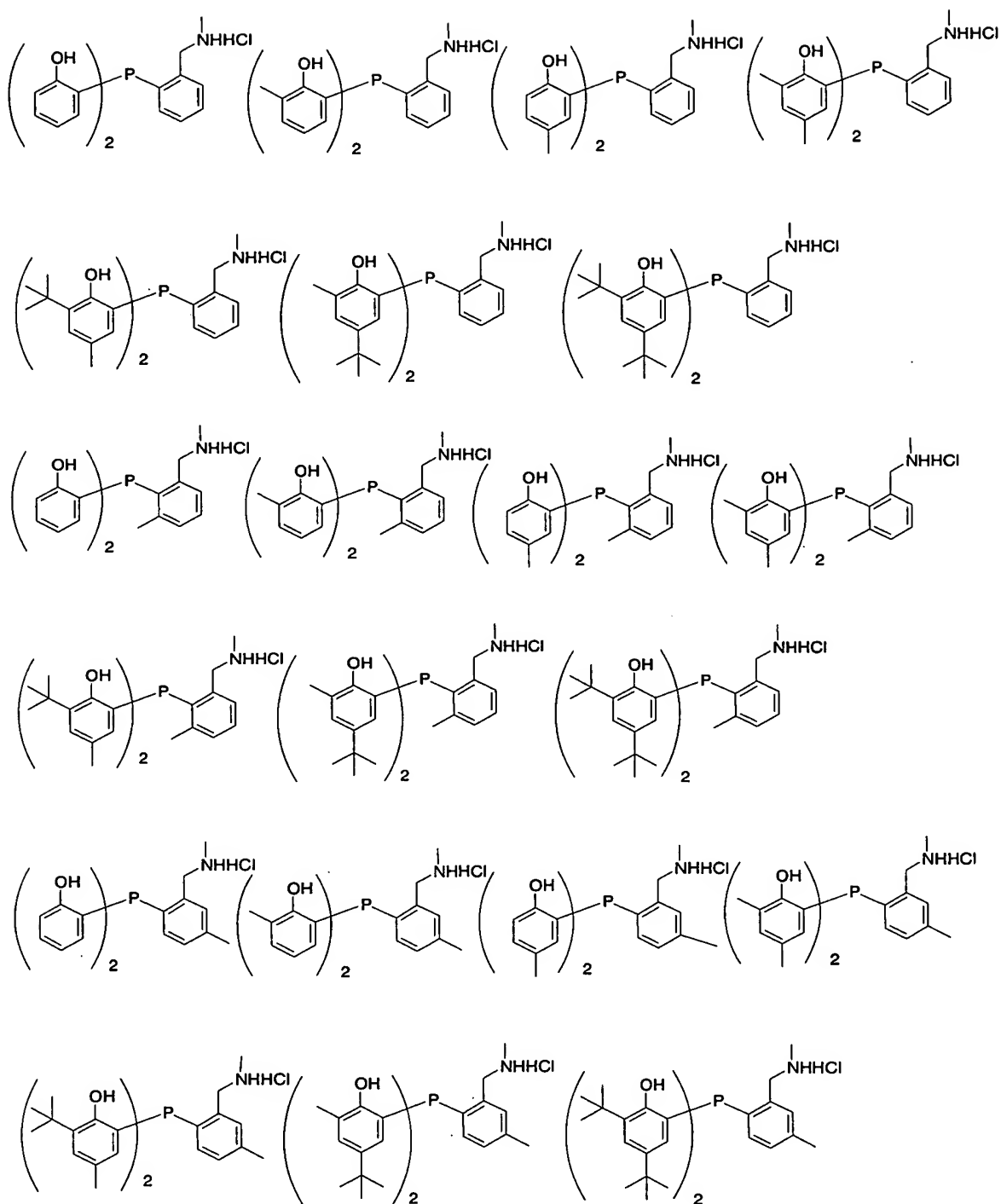


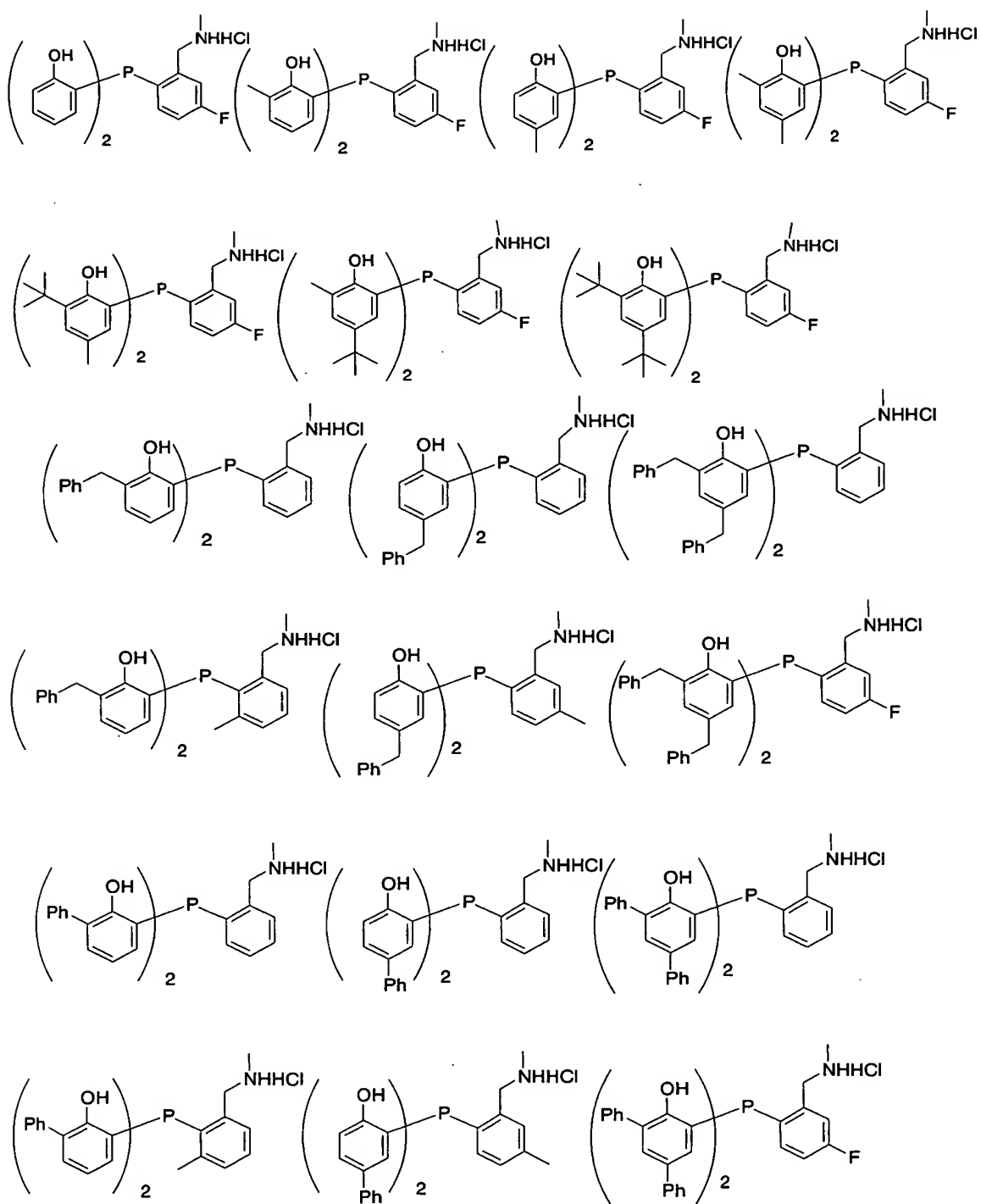


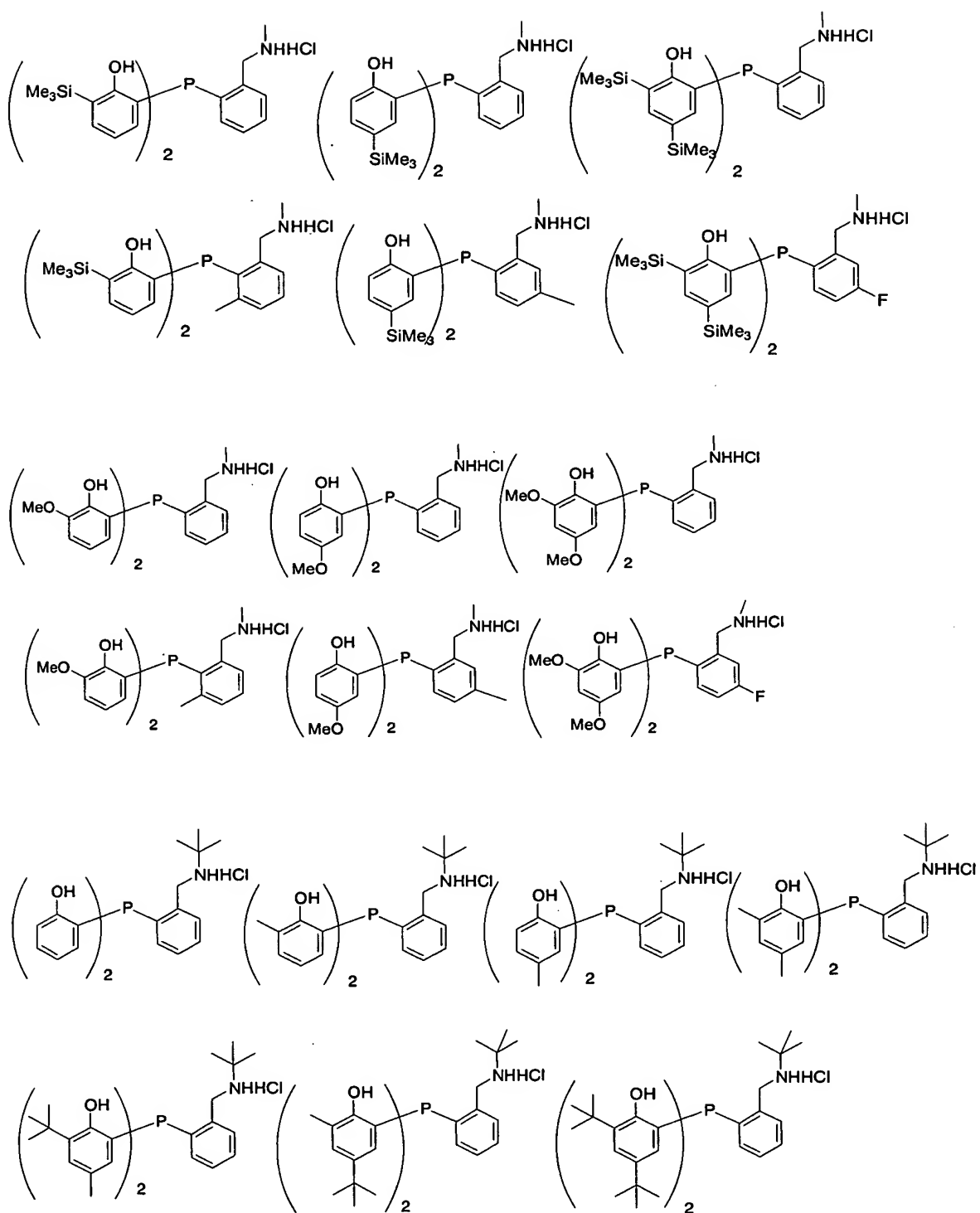


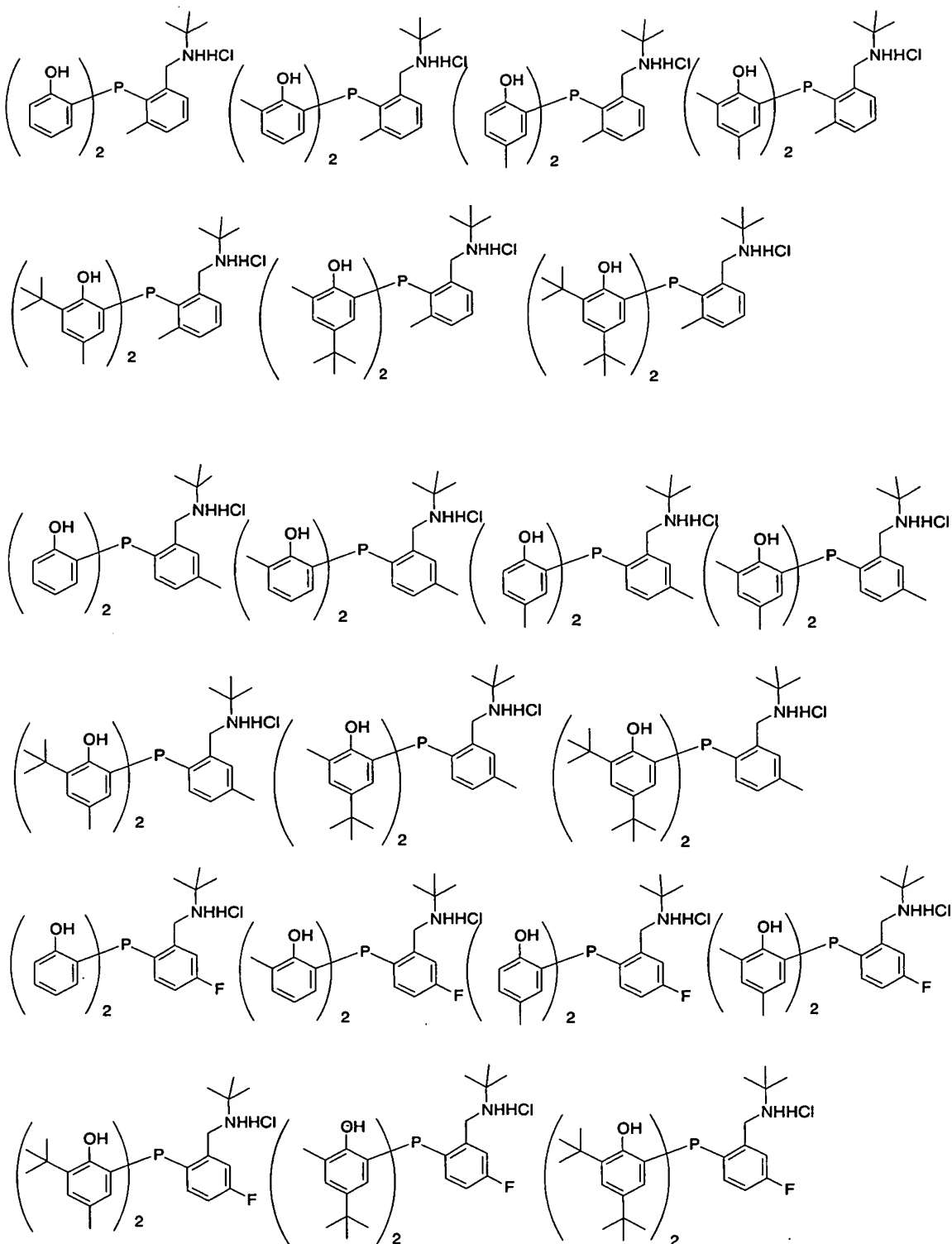


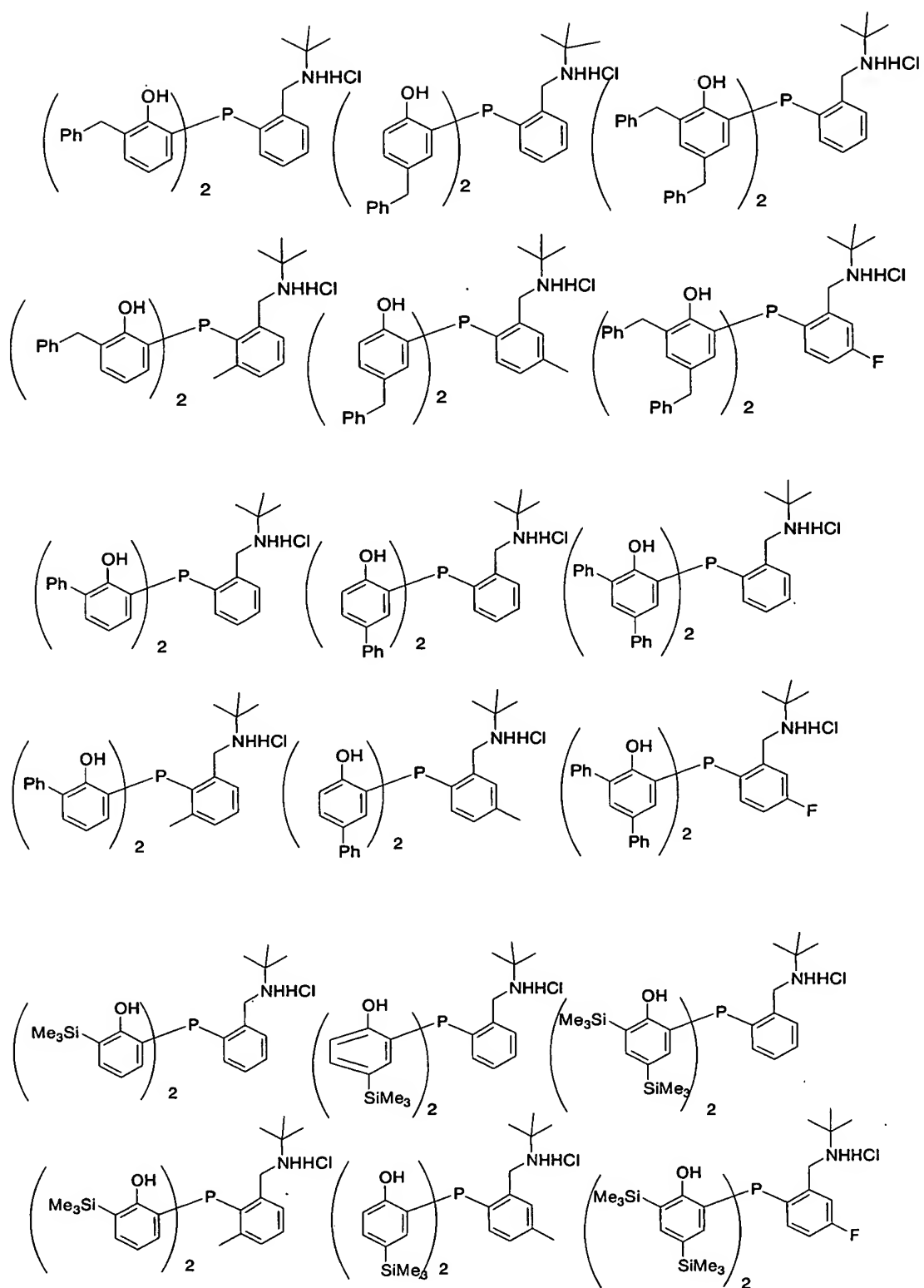
Specific examples of hydrogen halide acid salts of the compound of formula (23A) include, for example, the following
 5 compounds:

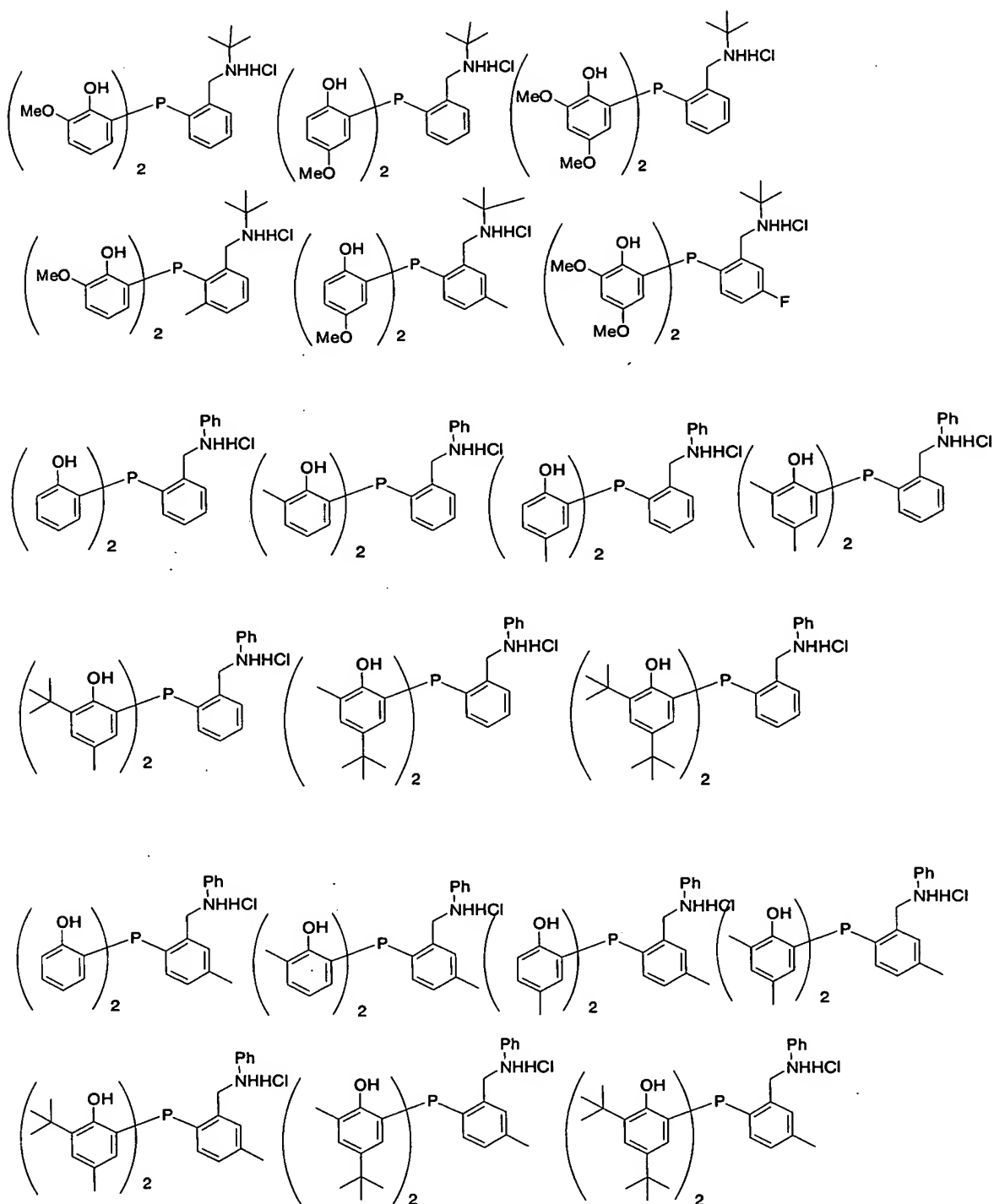


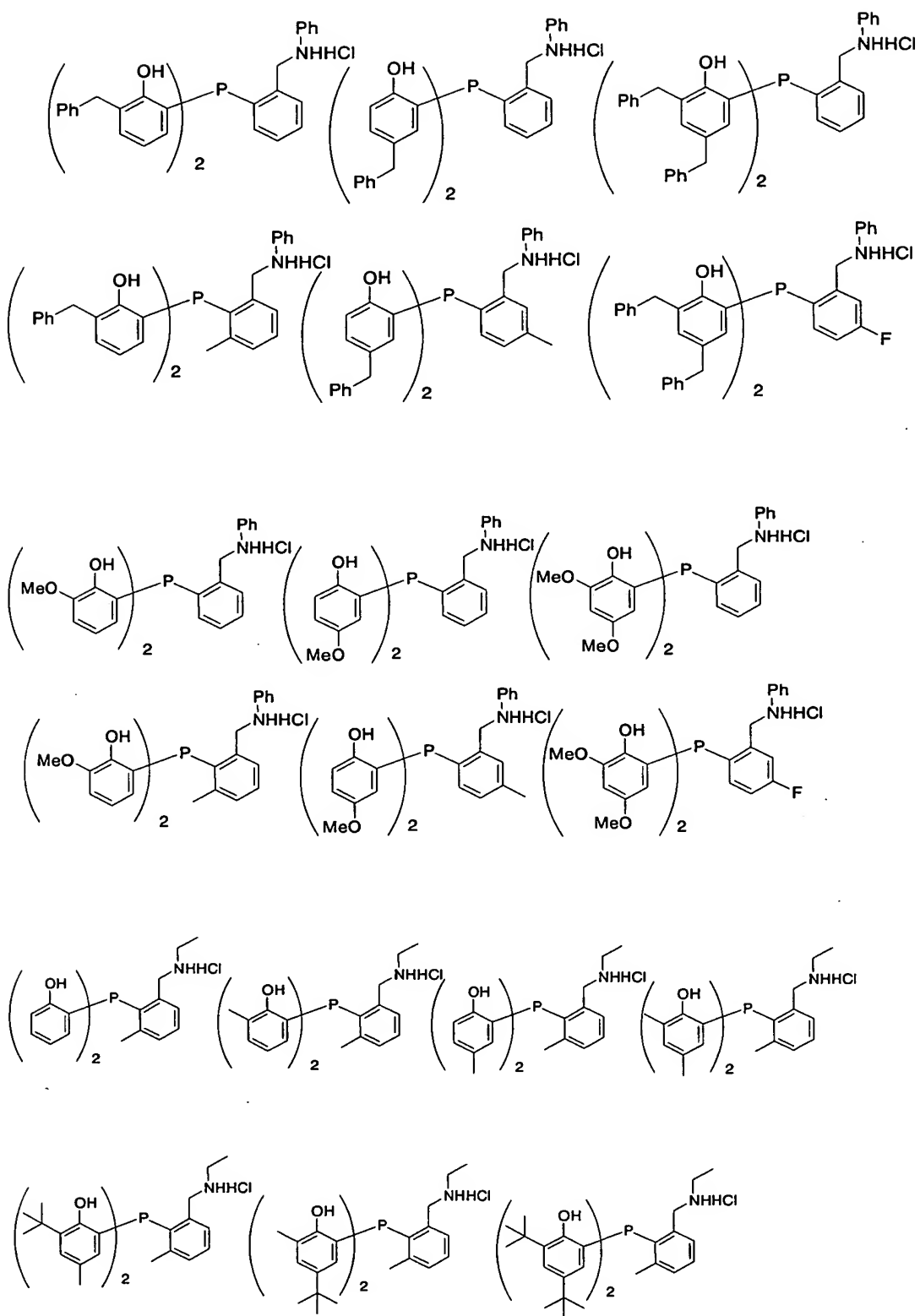


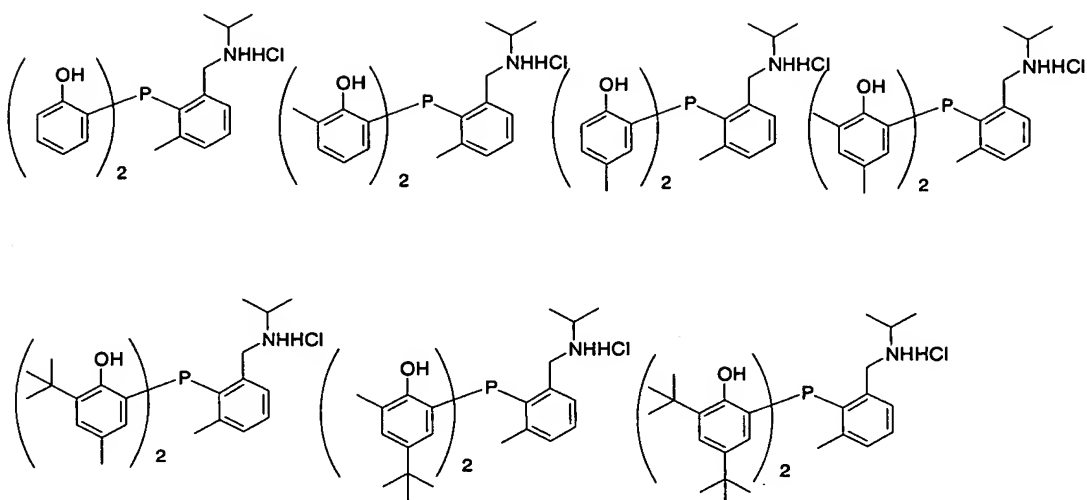




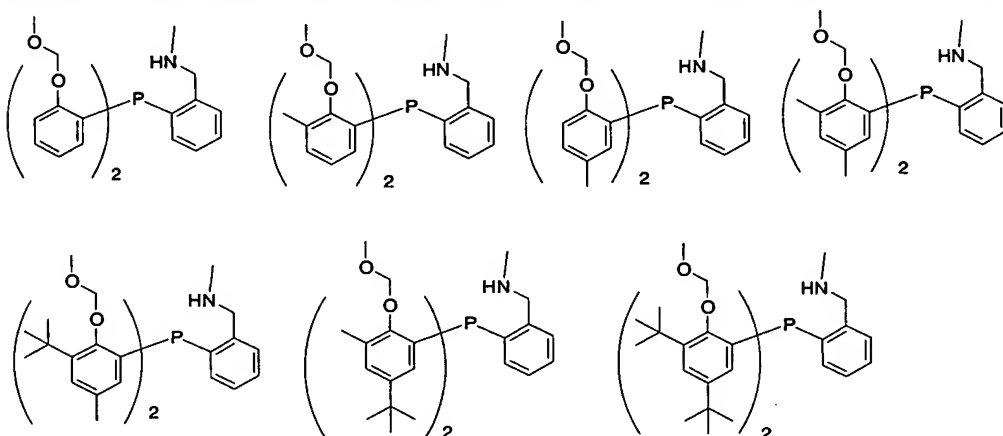


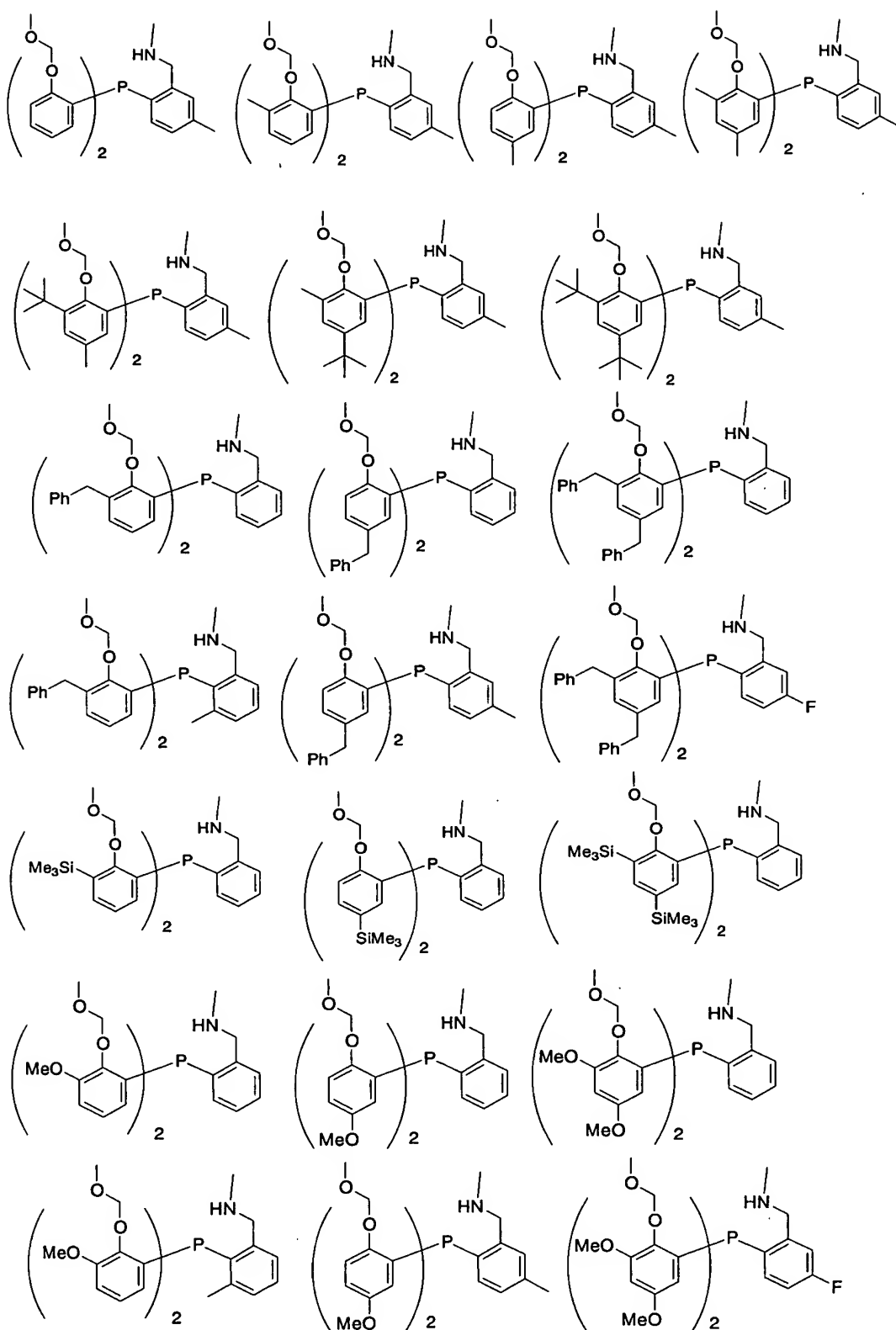


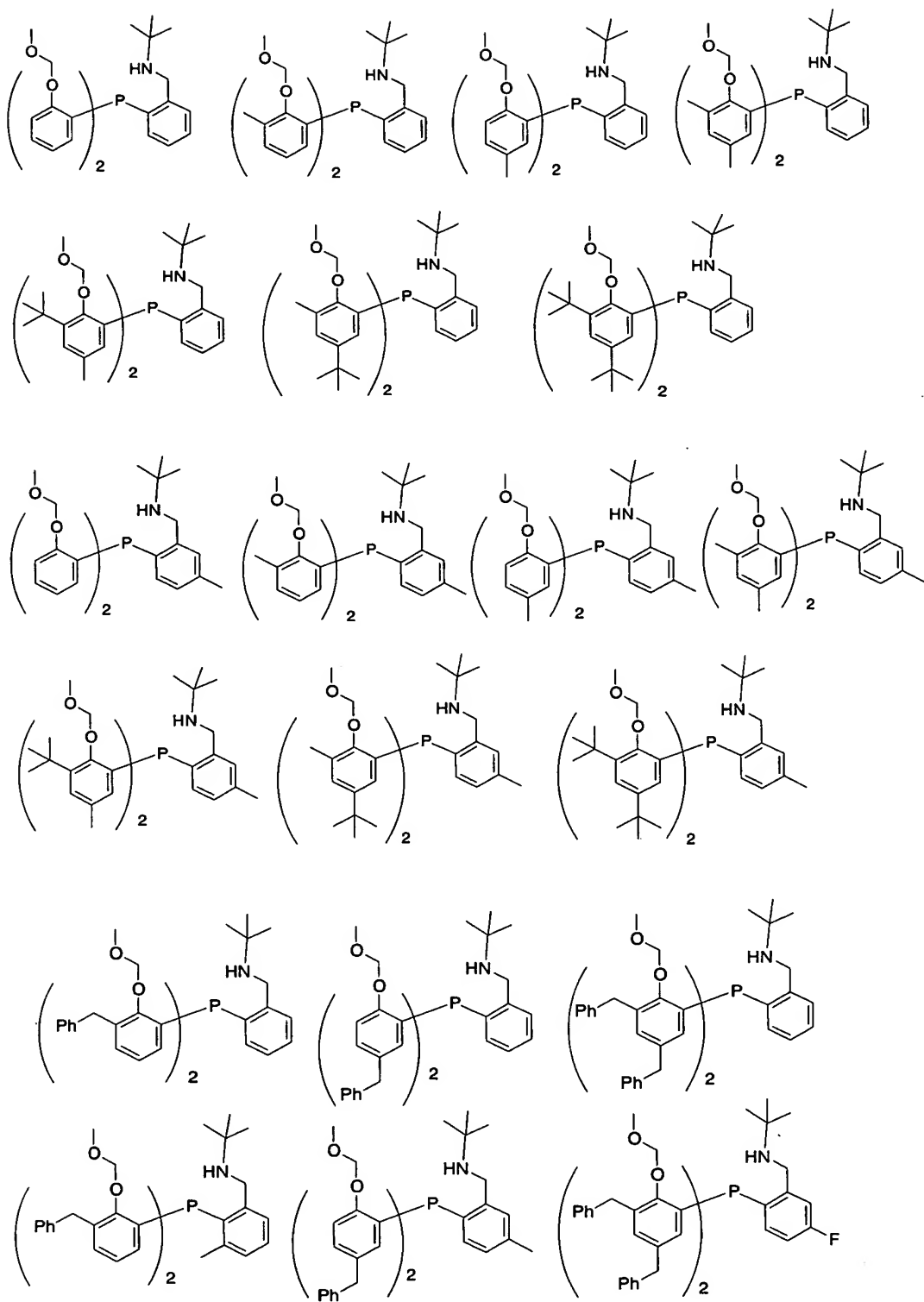




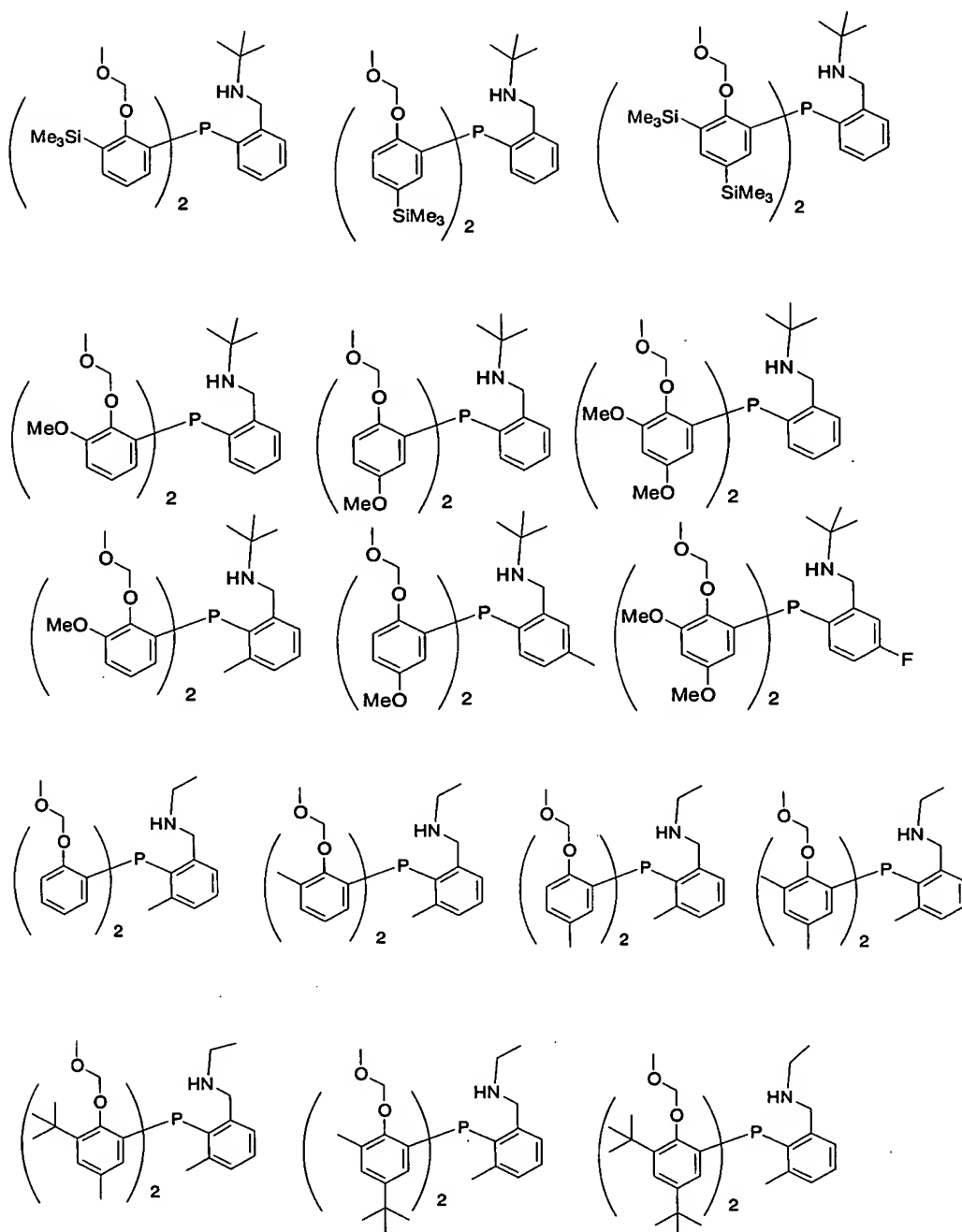
Specific examples of the phosphine compound of formula (1) wherein G^1 is G^{23} , which corresponds to the compound of formula 5 (23B) include, for example, the following compounds:



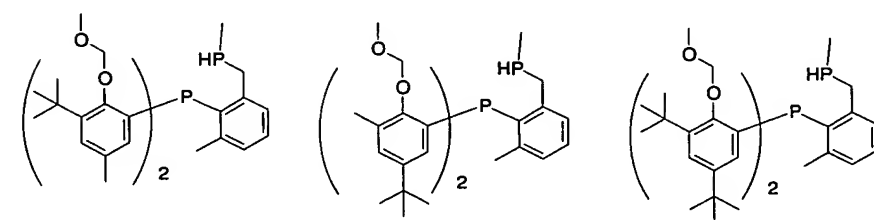


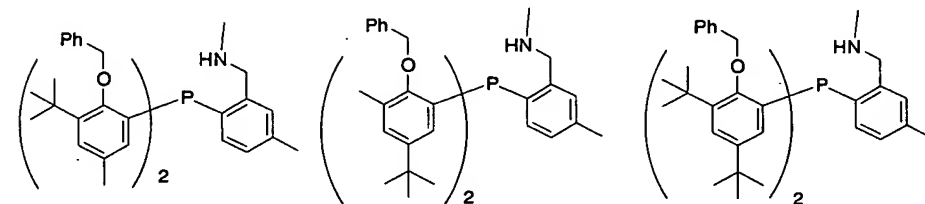
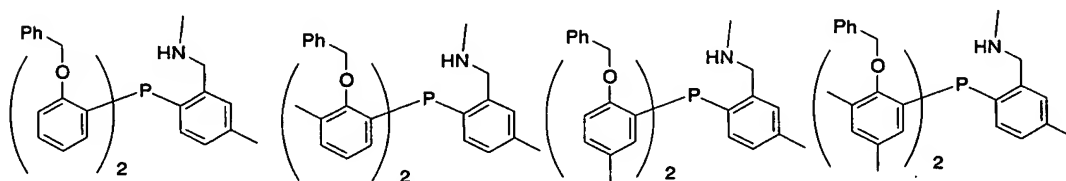
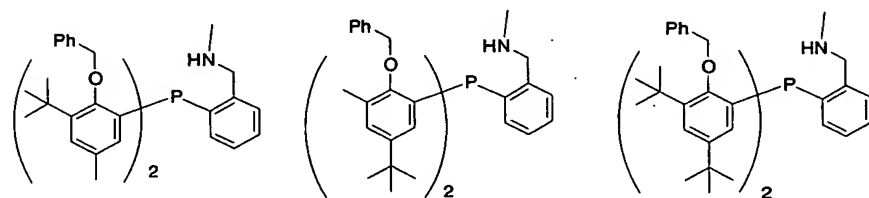
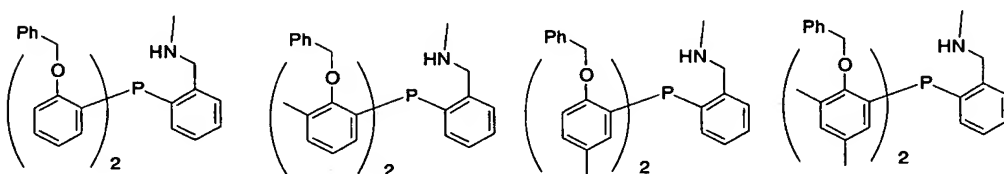
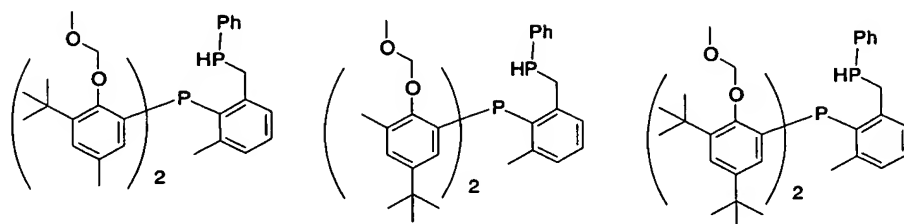
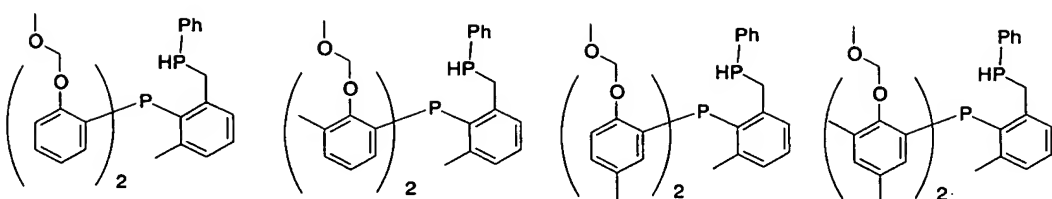


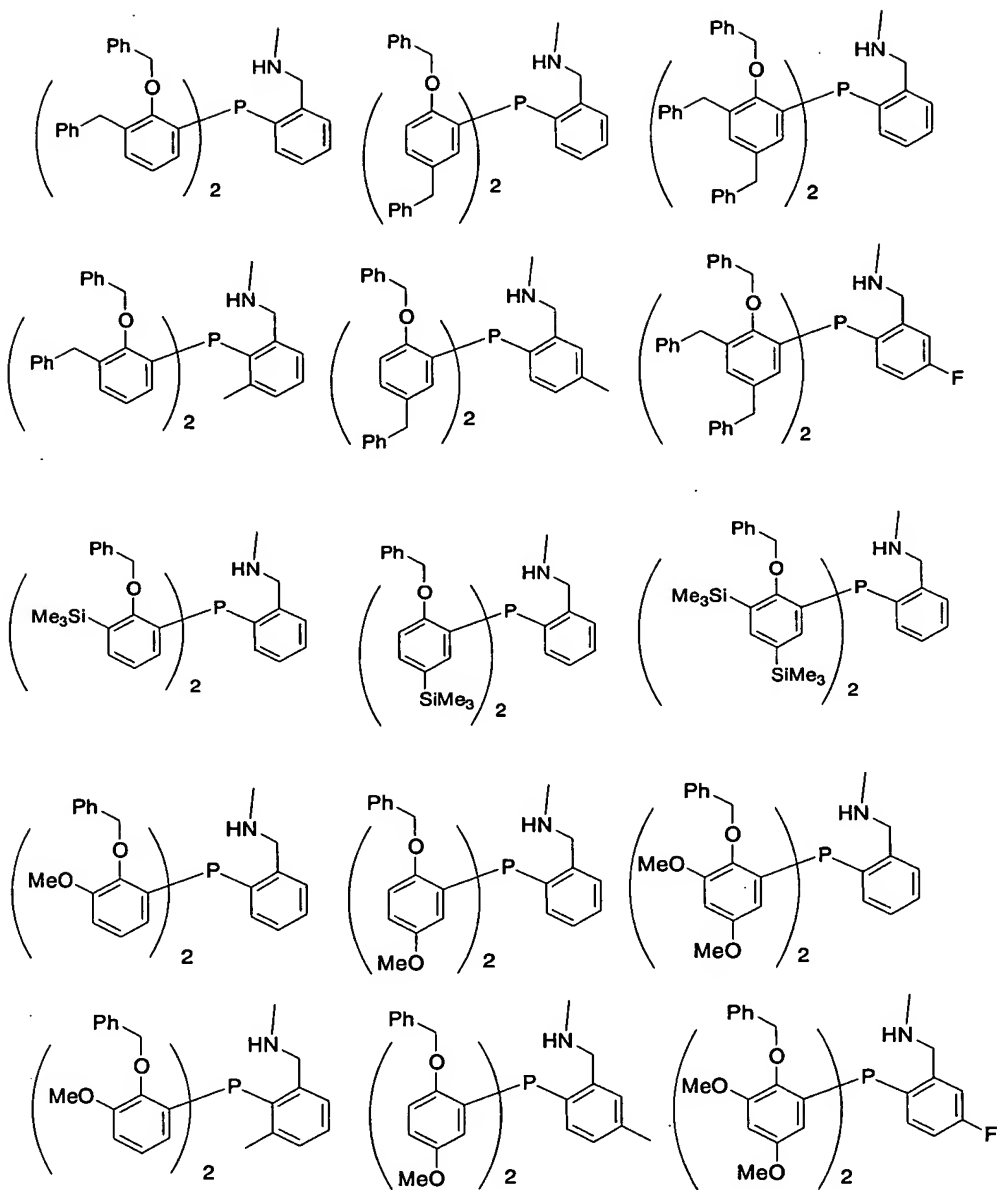
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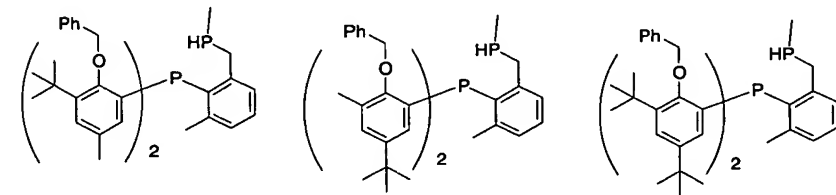
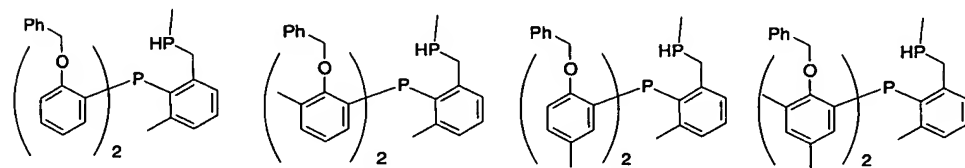
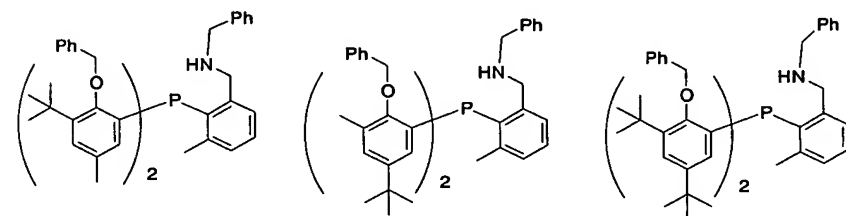
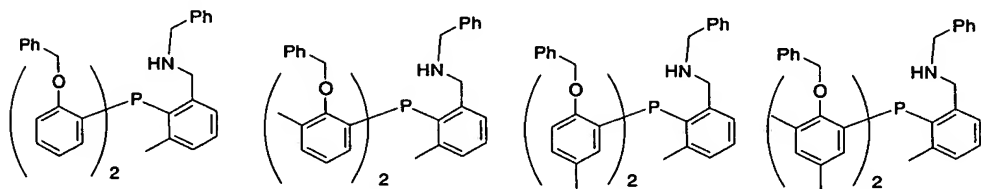
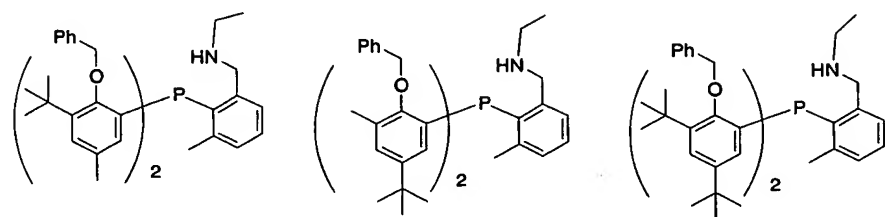
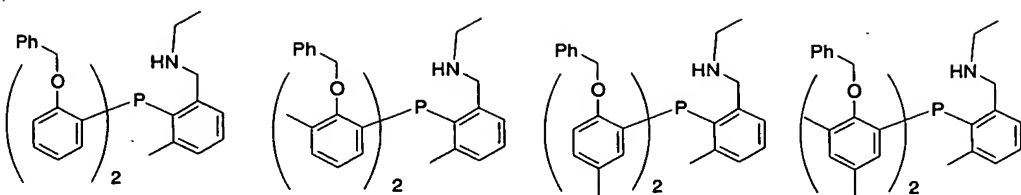
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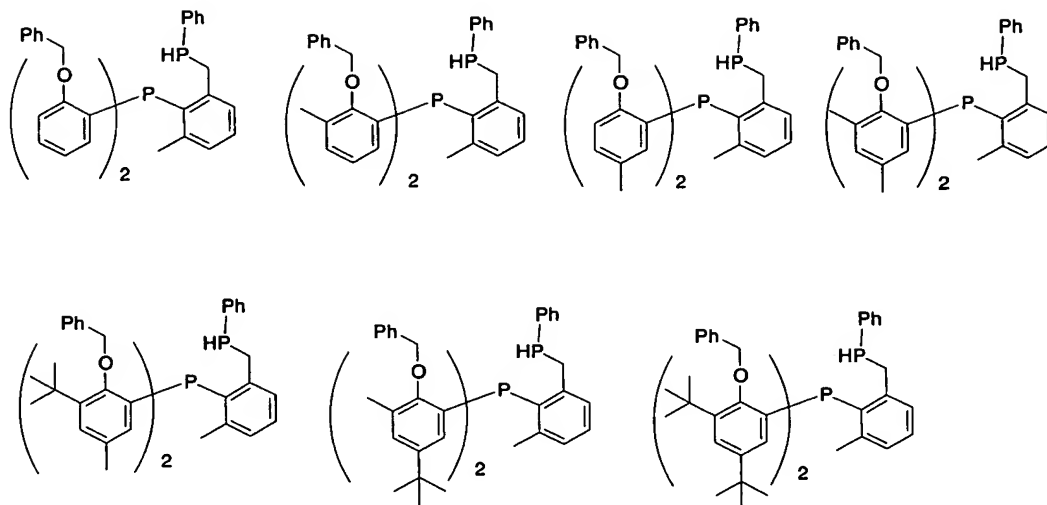






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The phosphine compound of formula (23B) where A¹ denotes a nitrogen atom can be produced by reducing the compound of formula (23C). The reducing reaction can be performed by using a metal hydride compound such as sodium borohydride, lithium aluminum hydride or the like, or hydrogen and the like.

The molar ratio between the compound of formula (23C) and metal hydride compound or hydrogen that may be used in the reaction is not particularly restricted, and it is preferably in the range of 1:0.1 to 1:10, more preferably in the range of 1:0.5 to 1:5.

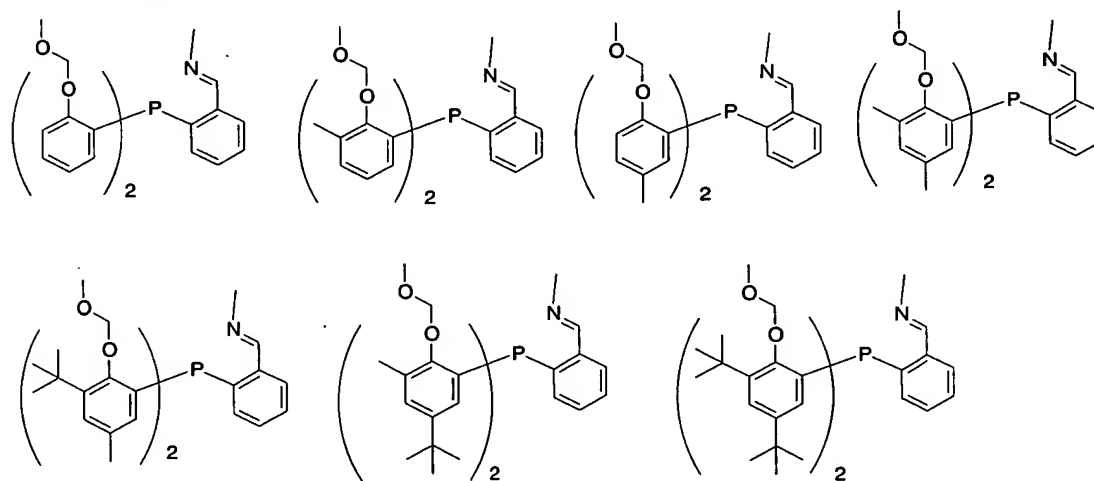
The reaction is usually performed in an organic solvent. Examples of the solvent include aprotic solvents including aromatic hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; ether solvents such as diethyl ether, tetrahydrofuran or the like; and halogenated solvents such as dichloromethane, dichloroethane, chlorobenzene, dichlorobenzene or the like; and protonic solvents such as

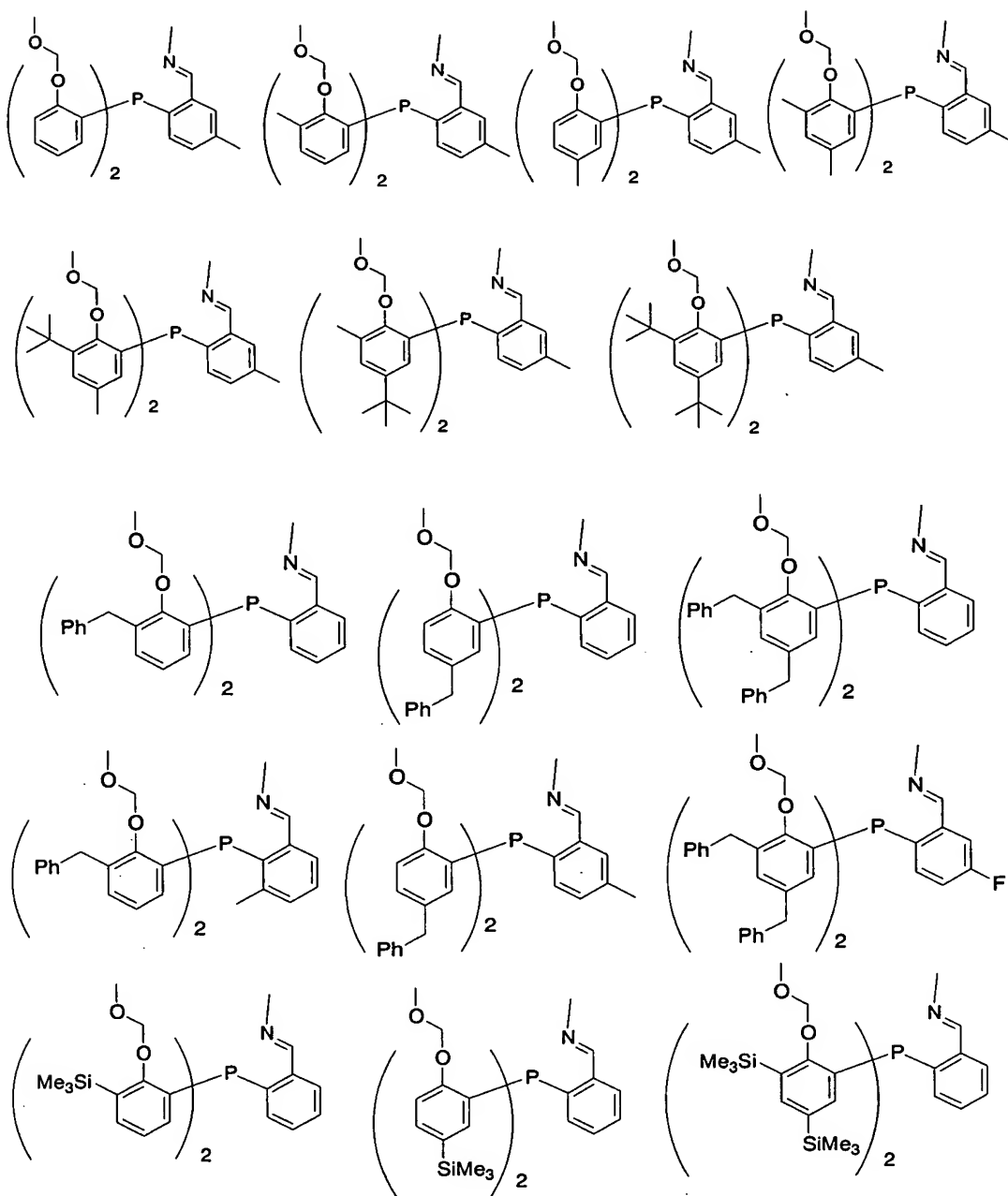
methanol, ethanol, isopropanol, butanol or the like. These solvents may be used alone or as a mixture of at least two of them, and the ratio thereof is usually in the range of 1 to 200 parts by weight, preferably in the range of 3 to 50 parts by weight, per part by weight of the triaryl compound of formula (23C).

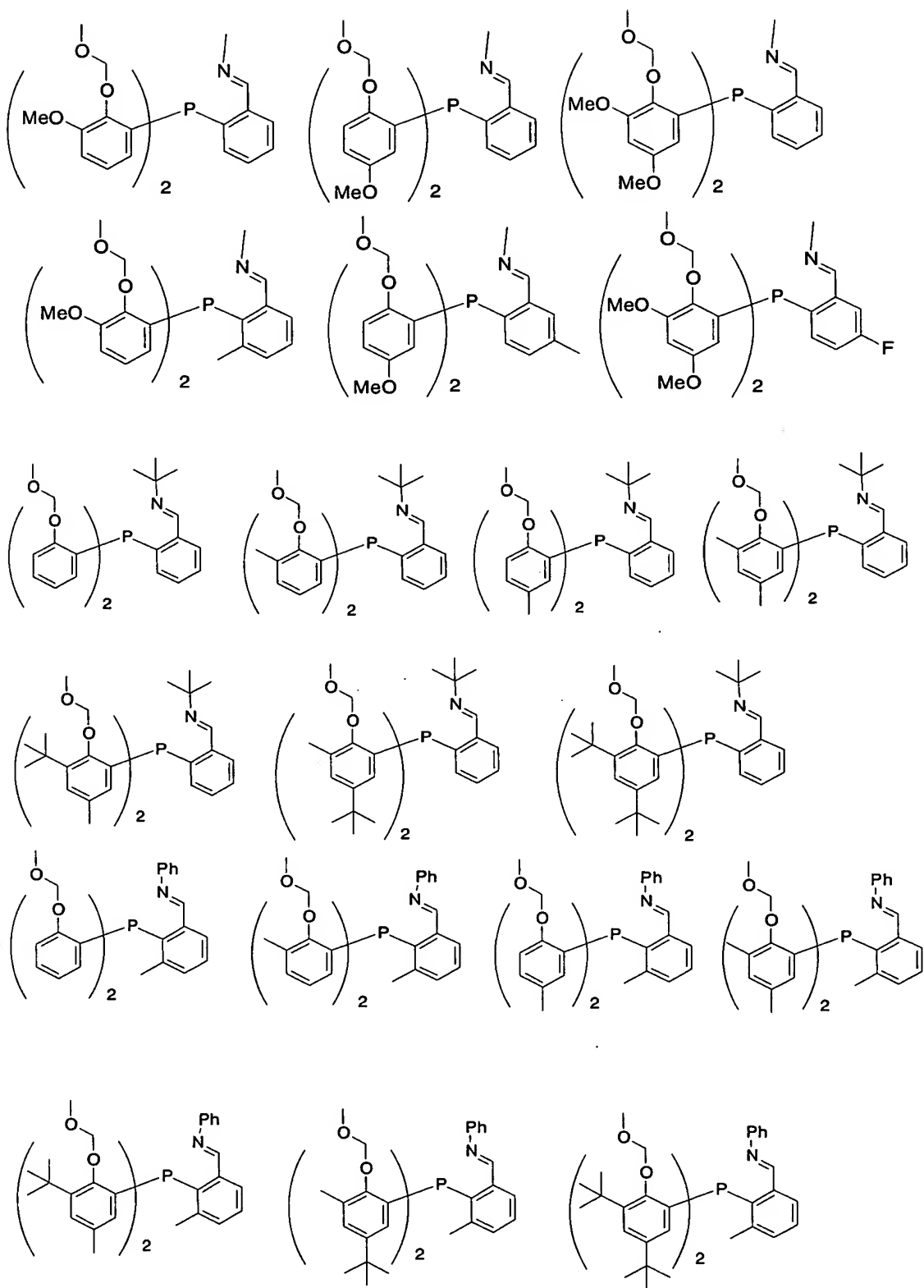
The reaction can be usually performed by adding the metal hydride compound or hydrogen to the compound of formula (23C). The reaction temperature is usually in the range from -100°C or more to the boiling point of the solvent or the less, more preferably in the range of -80°C to 100°C.

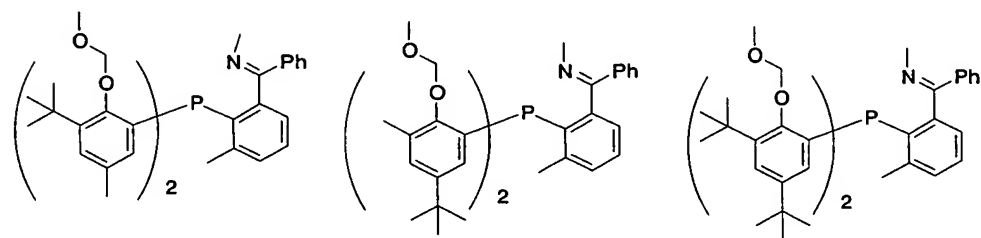
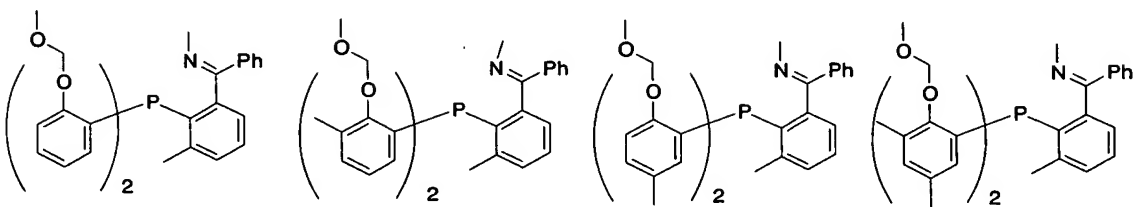
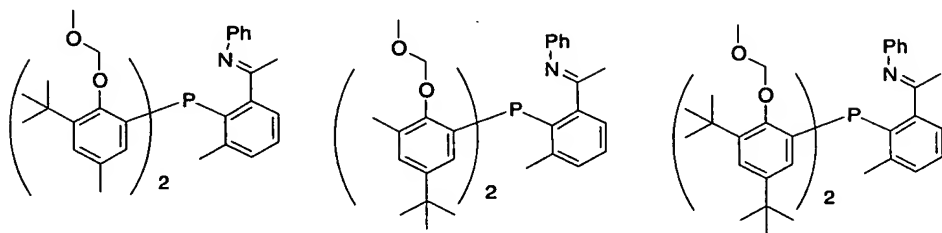
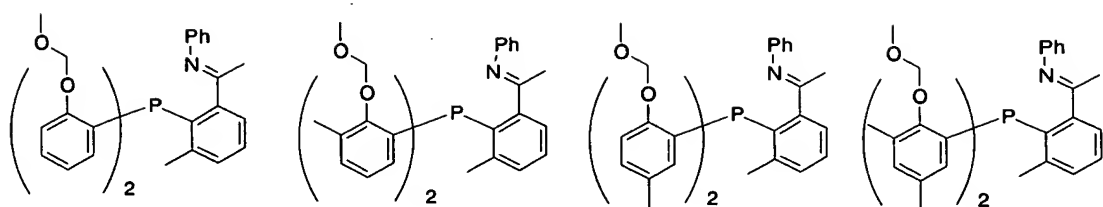
The phosphine compound of formula (23B) may be obtained from the reaction mixture obtained by removing the solvent by evaporation. Or, the compound may be purified by silica gel chromatography, if necessary.

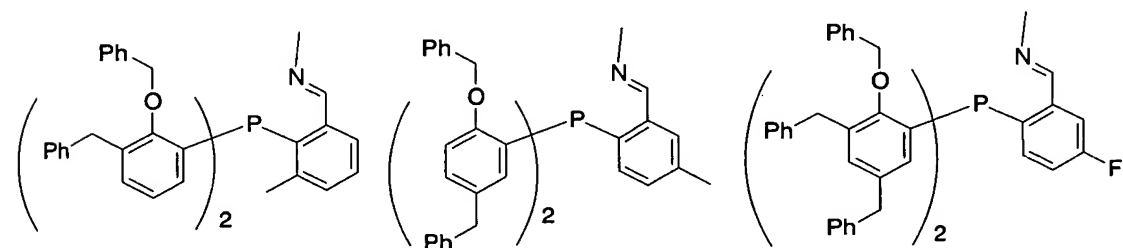
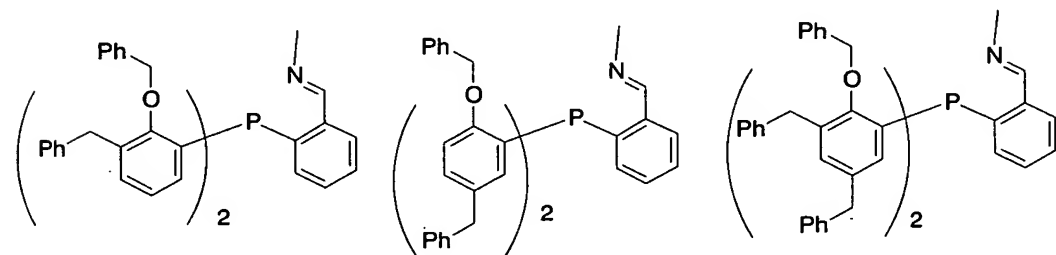
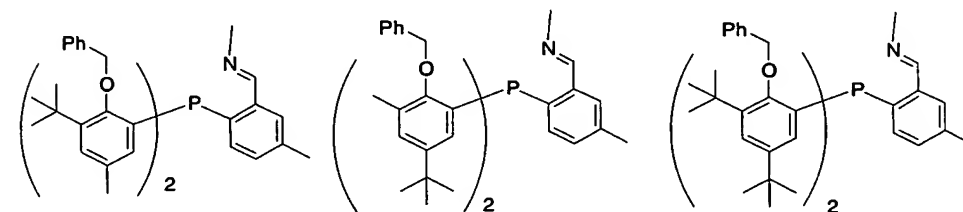
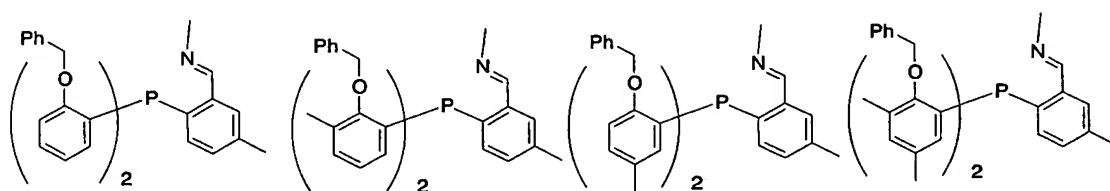
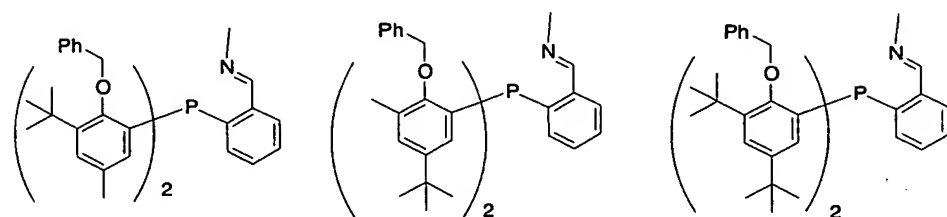
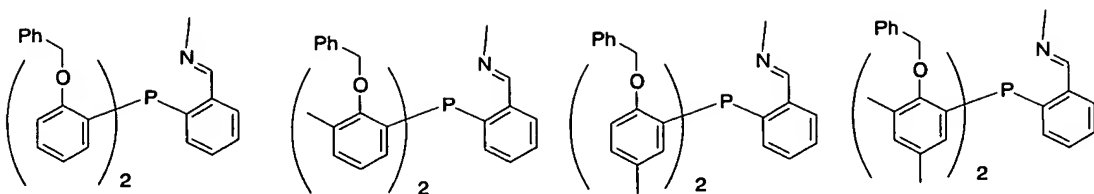
Examples of the compound of formula (23C) include the following compounds:

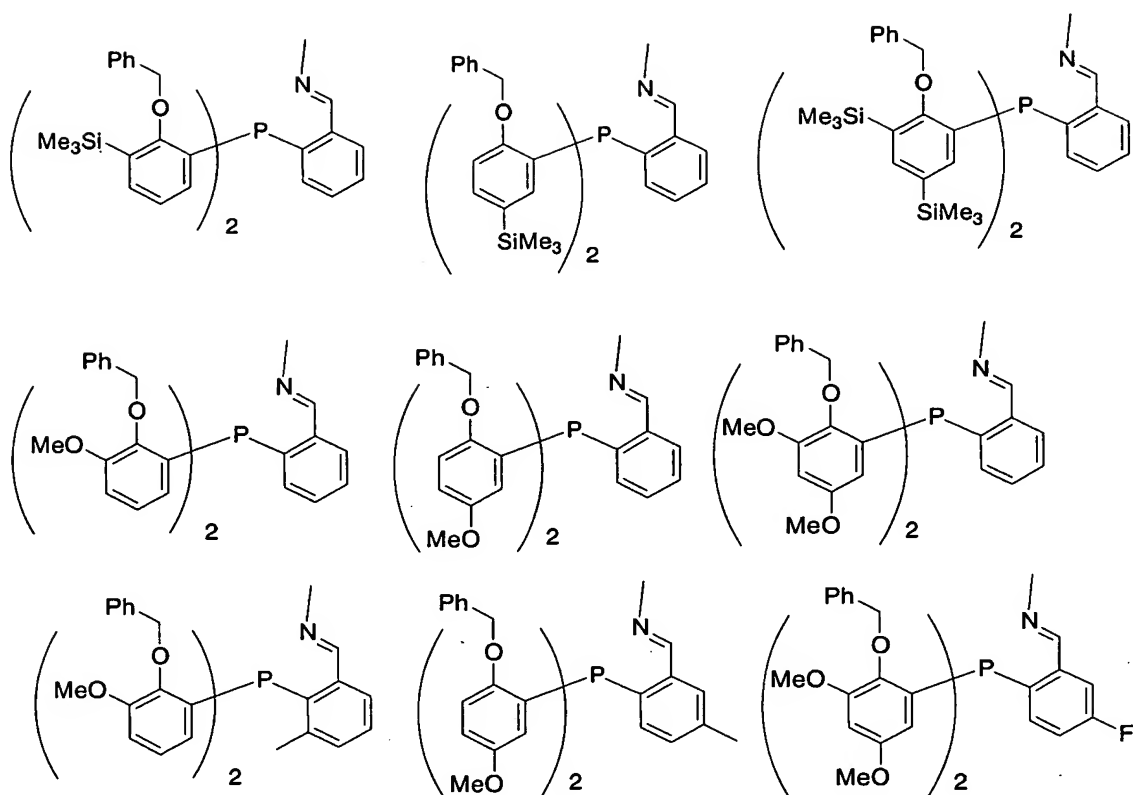




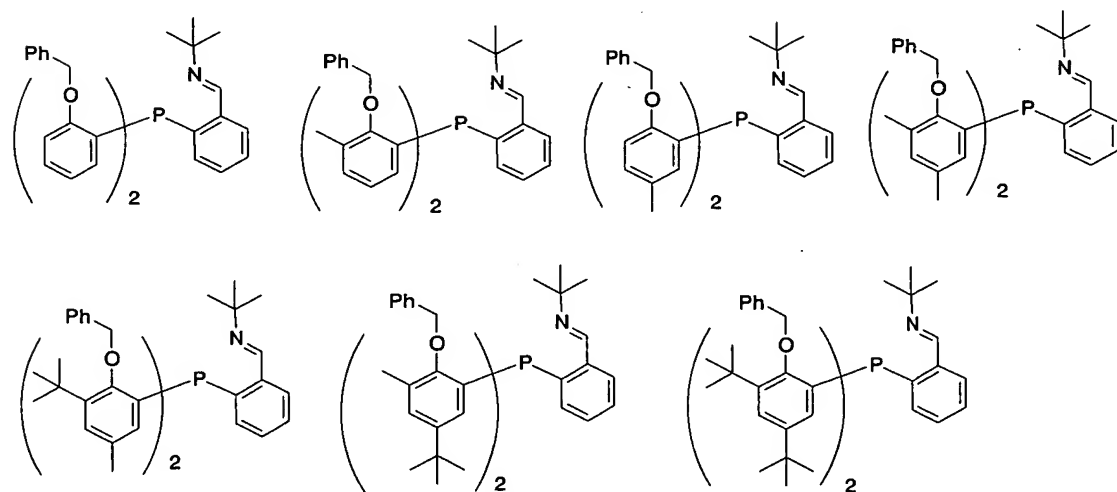


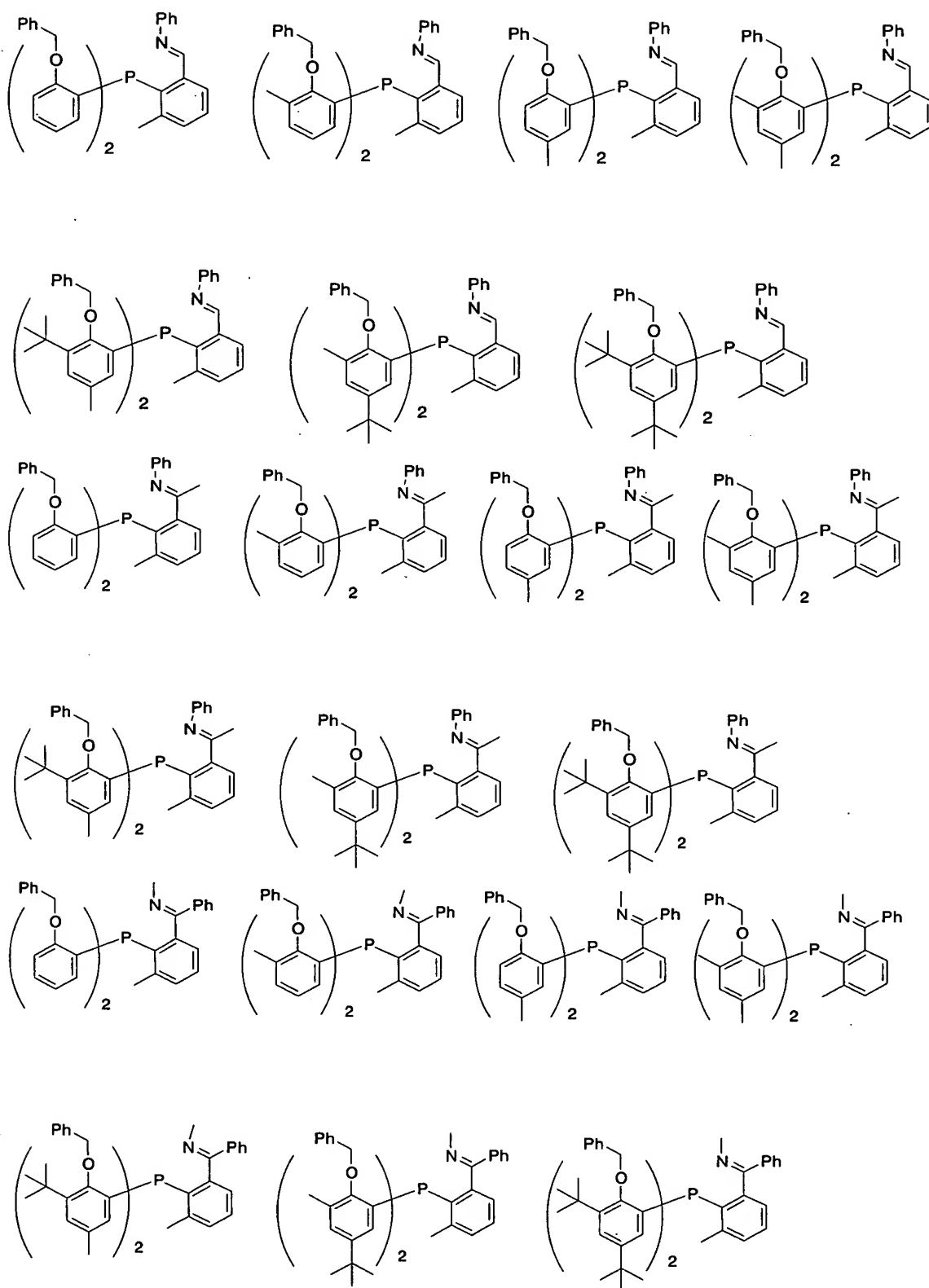




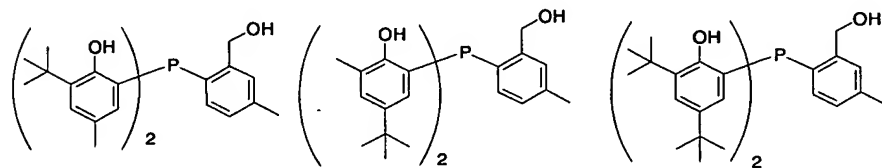
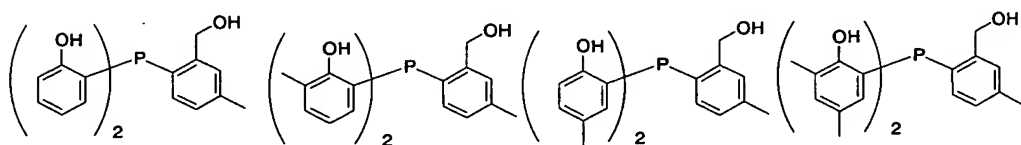
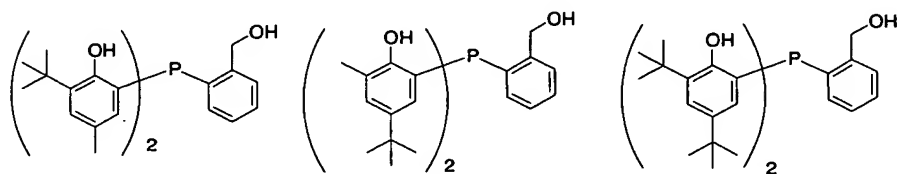
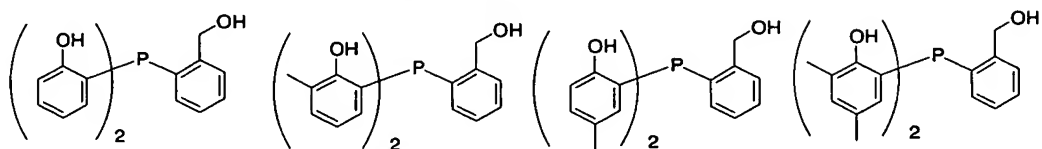


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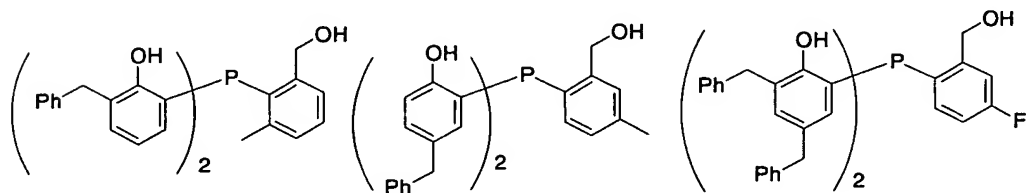
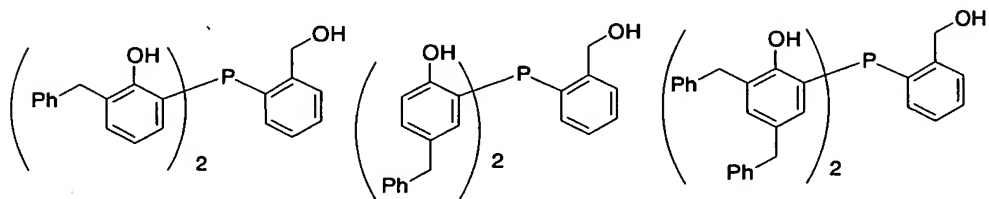


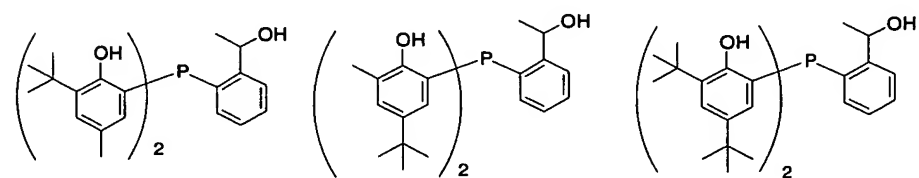
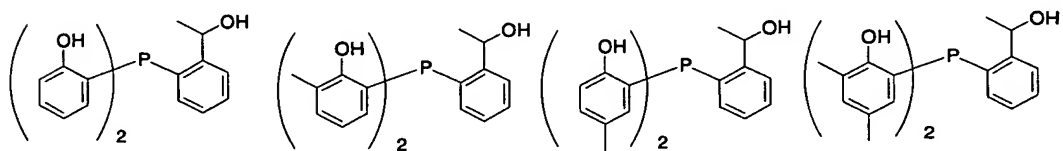
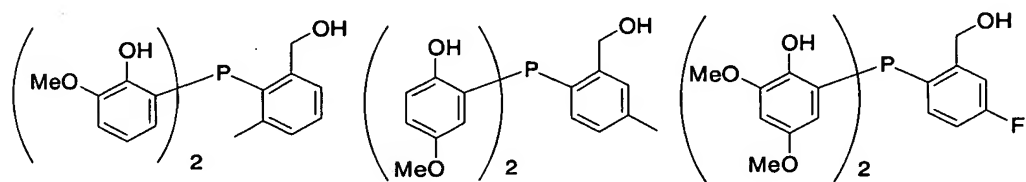
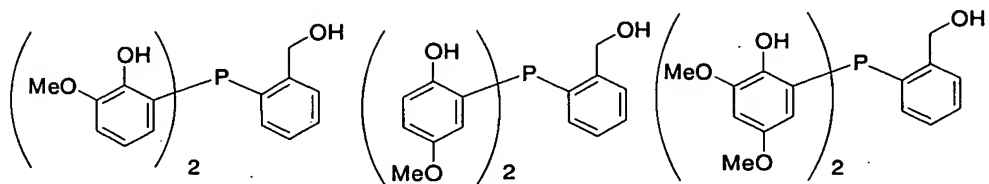
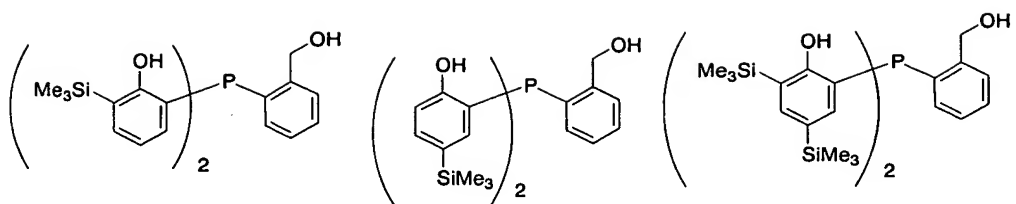


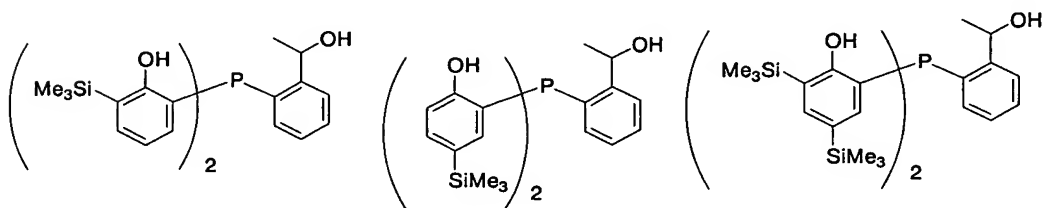
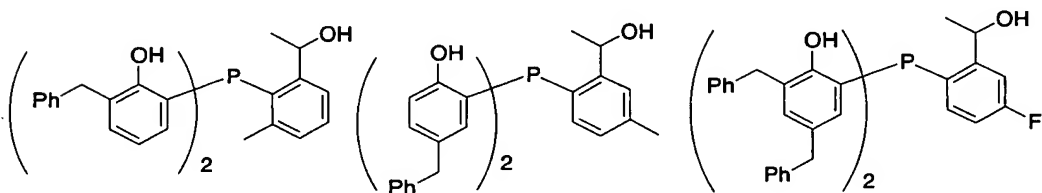
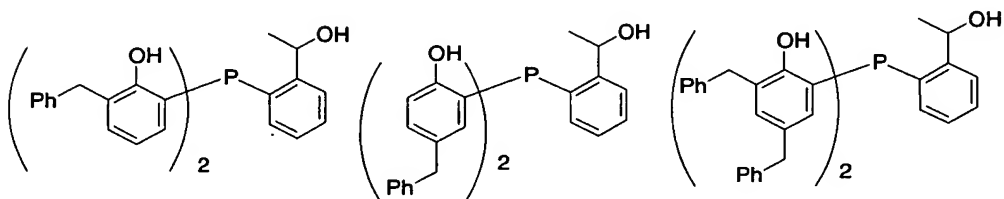
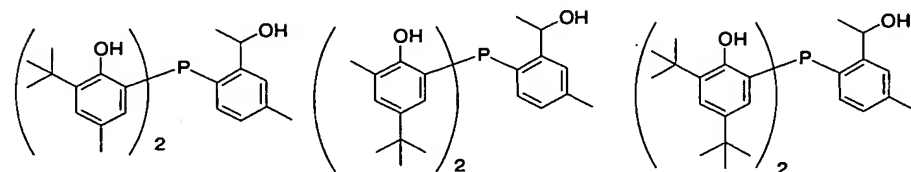
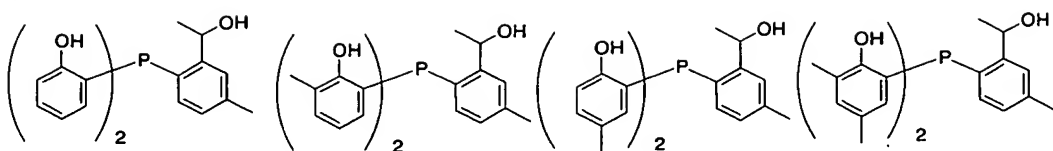
Examples of the compound of formula (1) wherein G^2 is G^{24} include the following compounds:



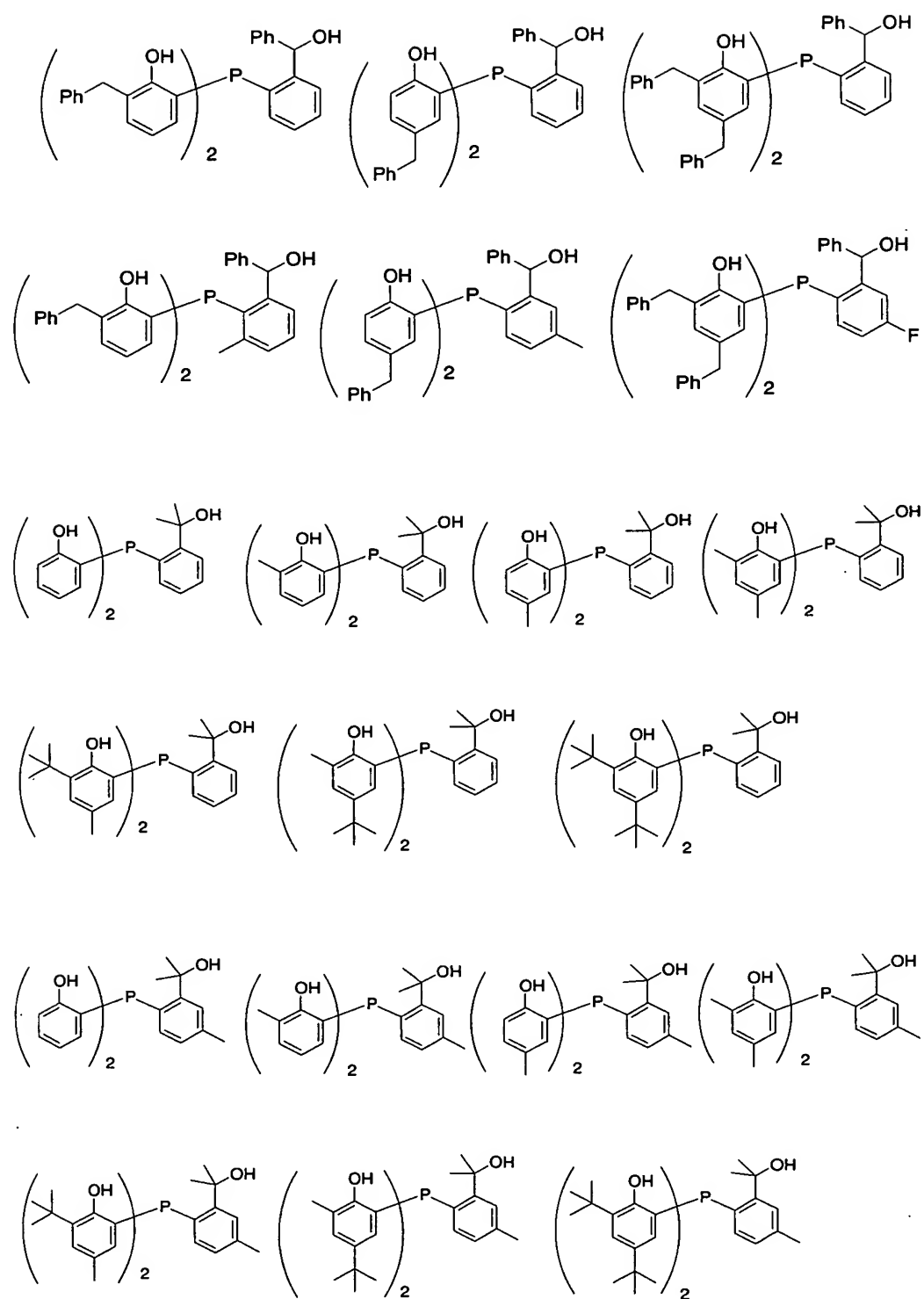
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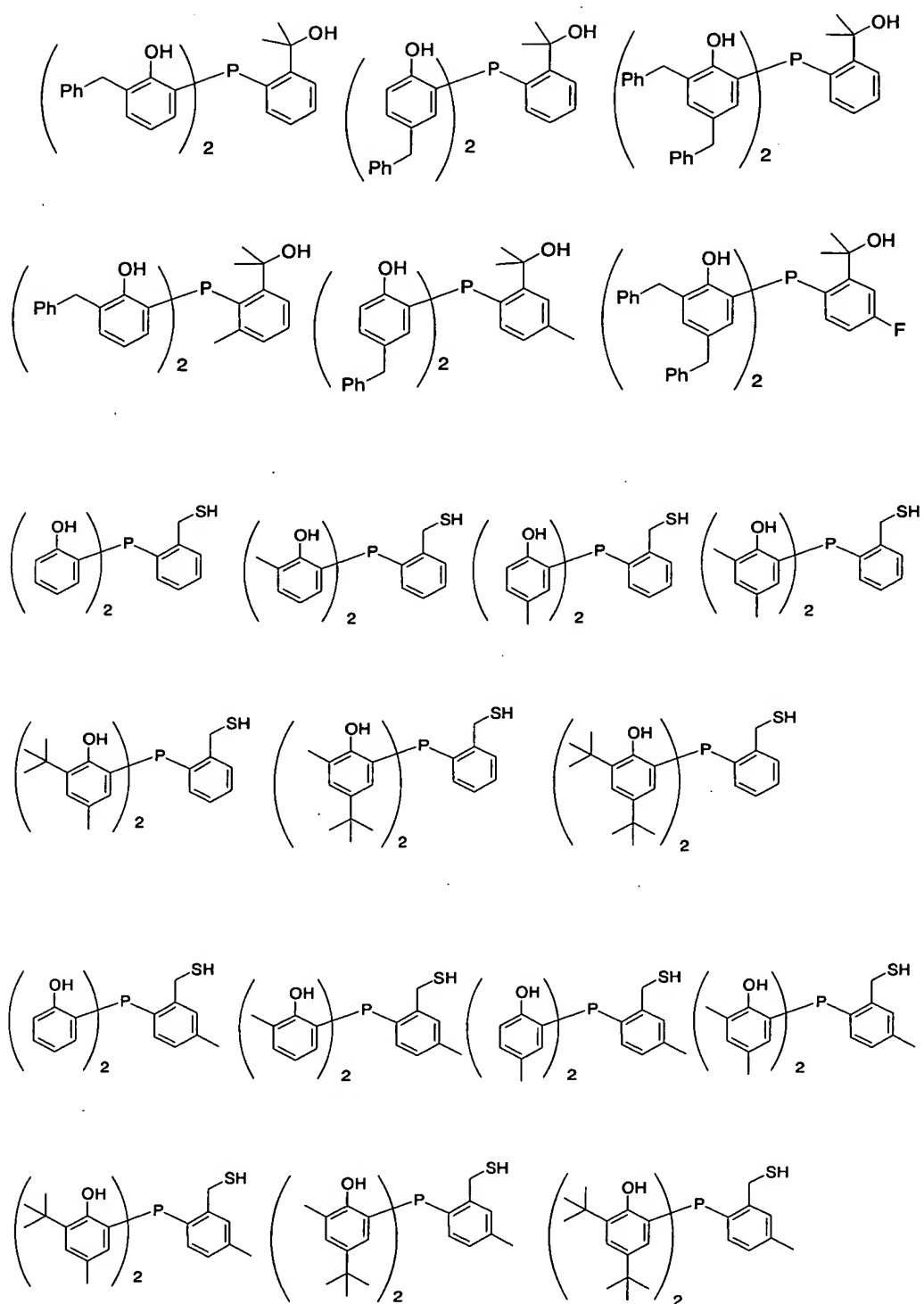


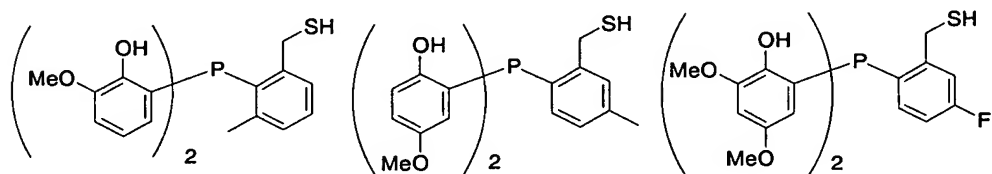
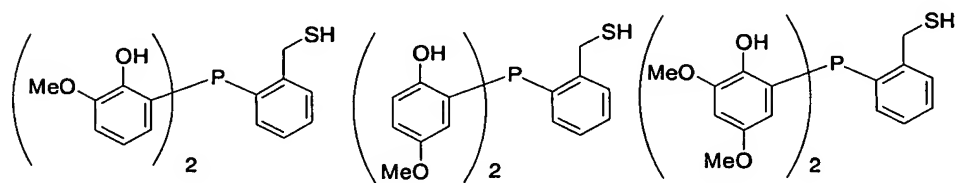
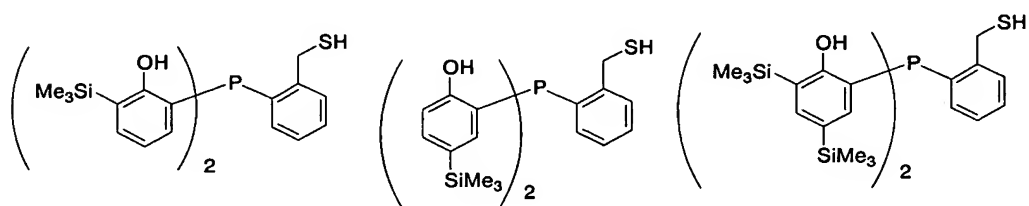
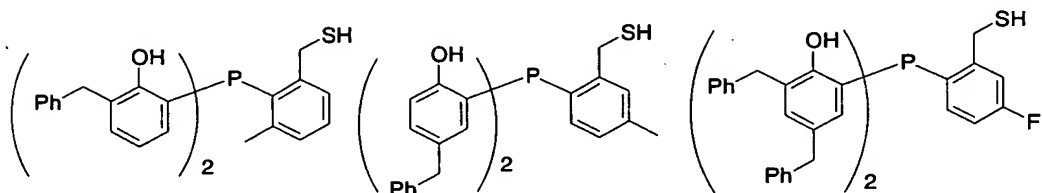
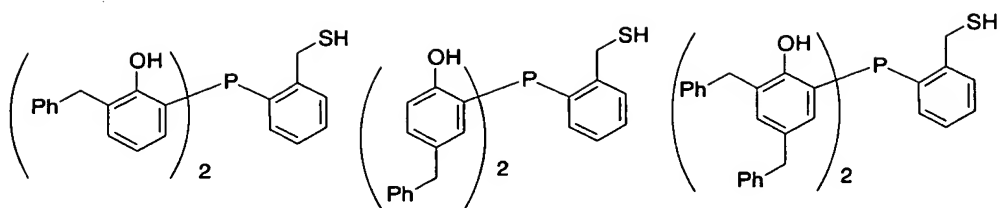




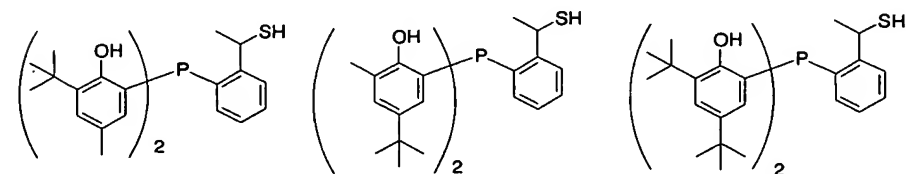
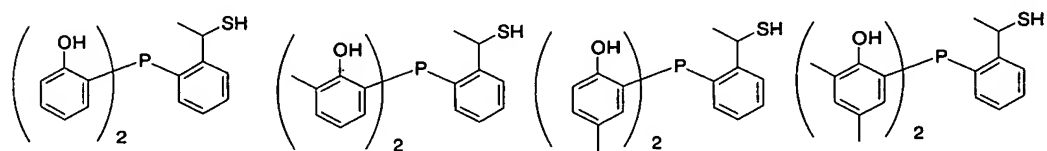


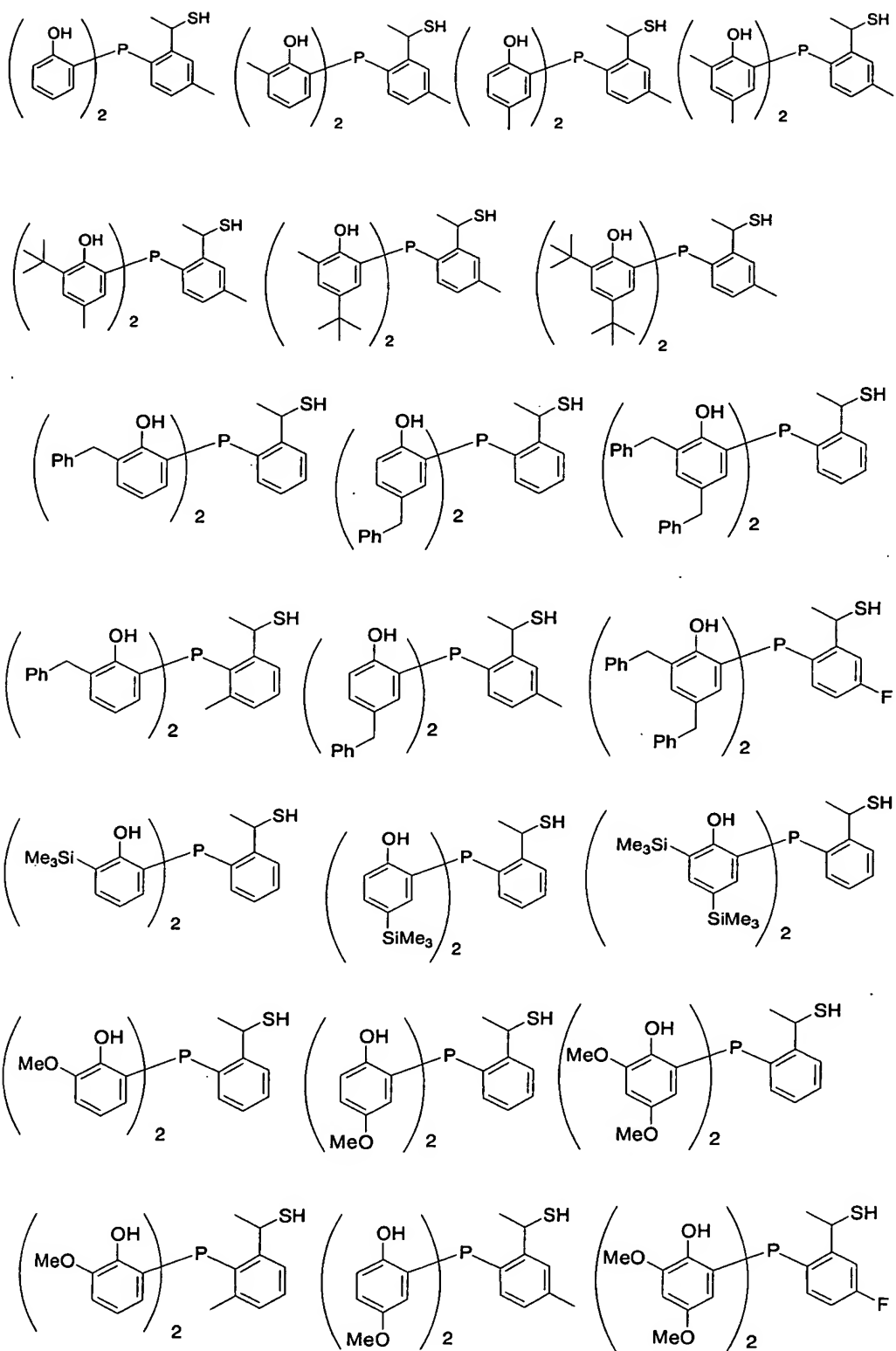






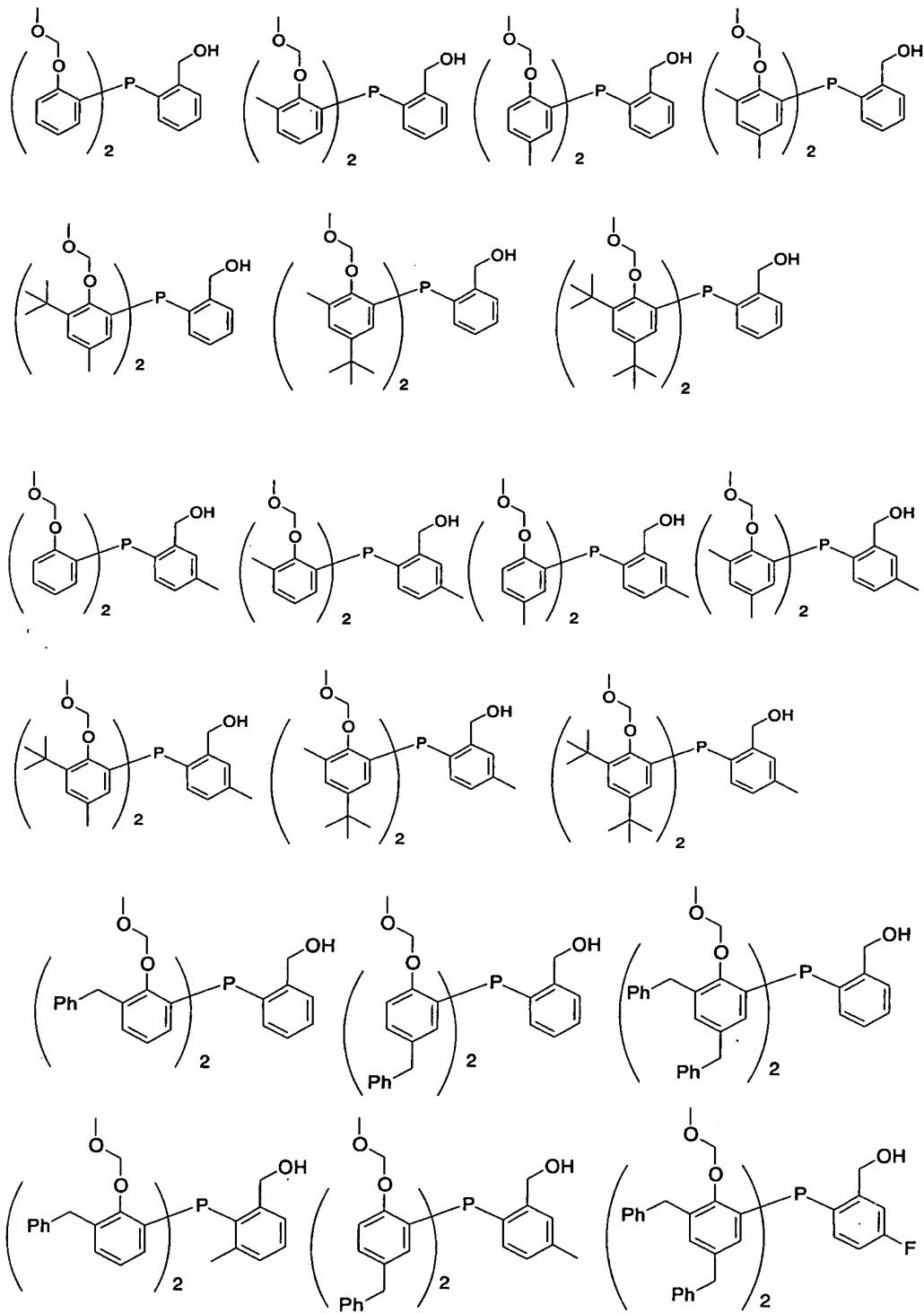
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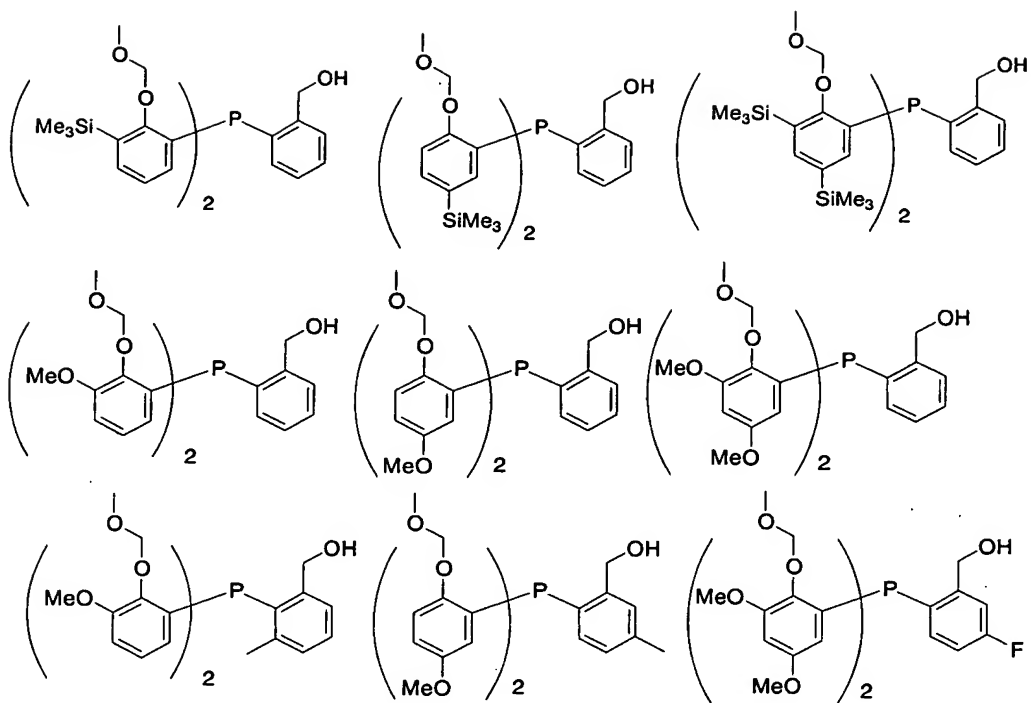




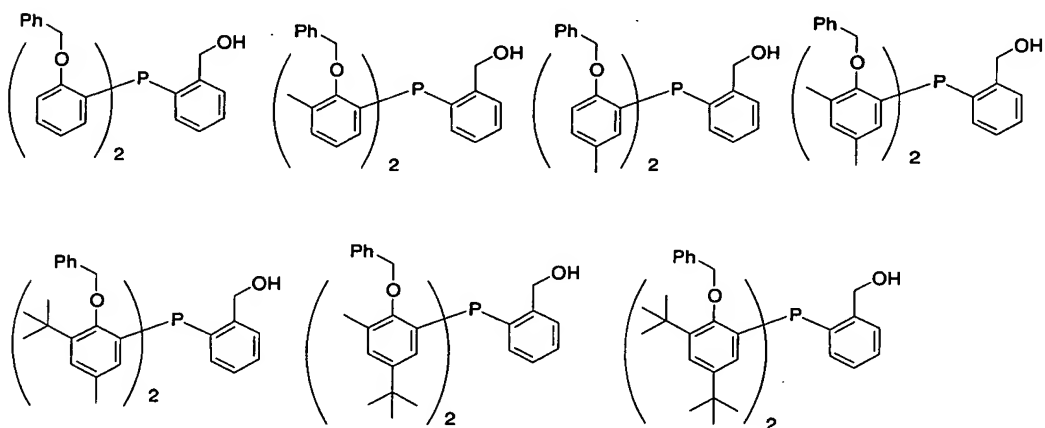
Examples of the phosphine compound of formula (1) wherein

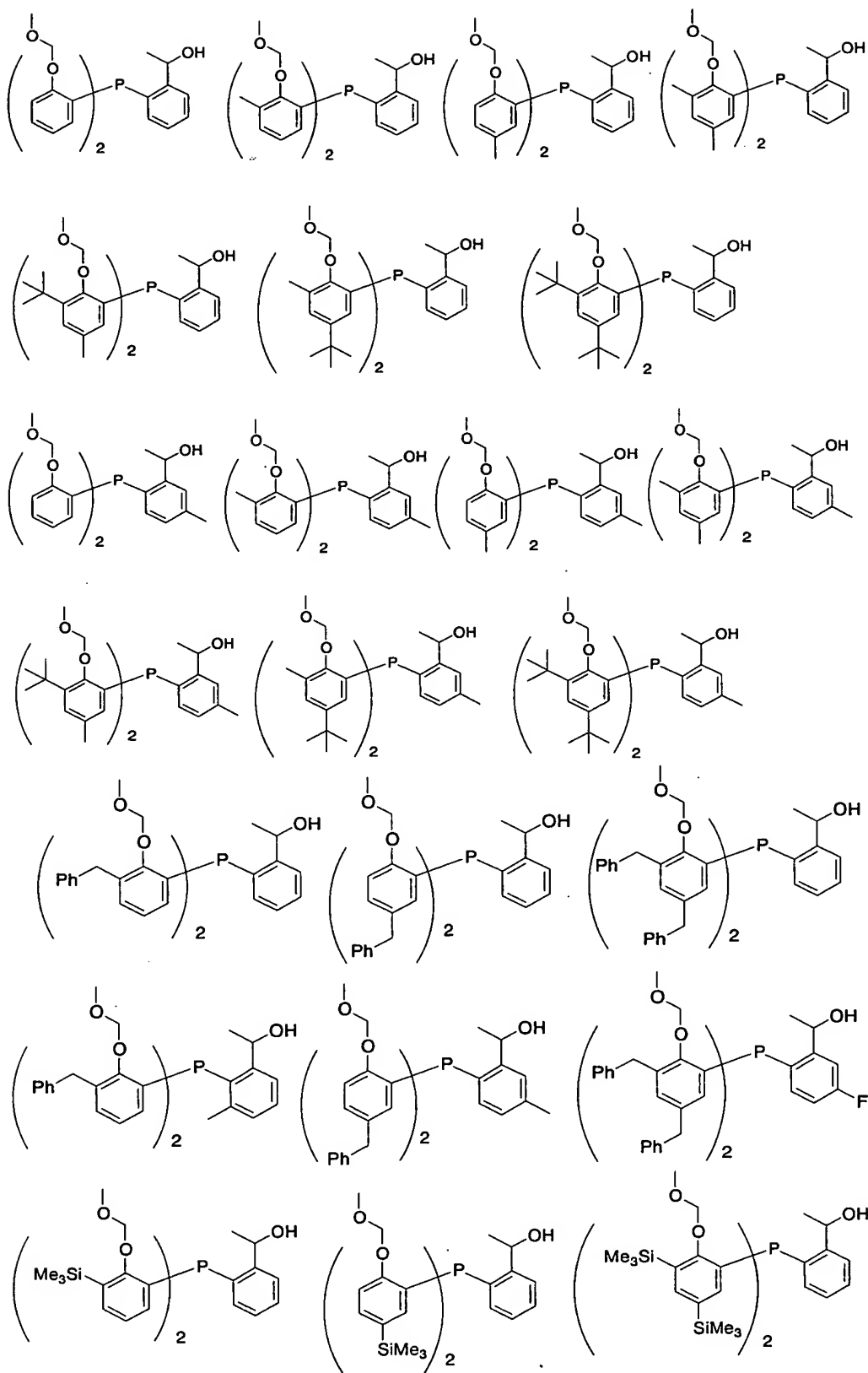
G^2 is G^{24} , or the compound of formula (24B), include, for example, the following compounds:

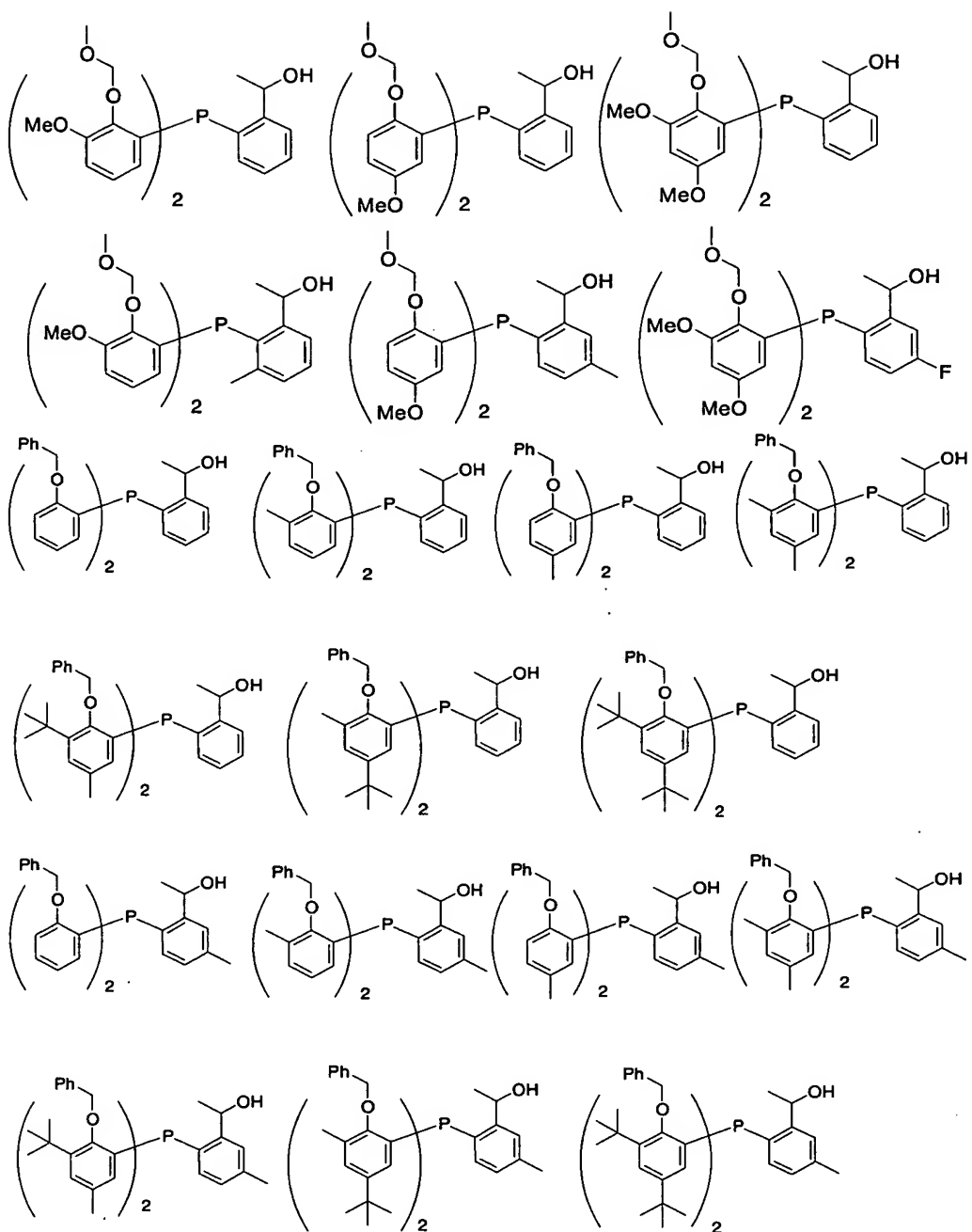


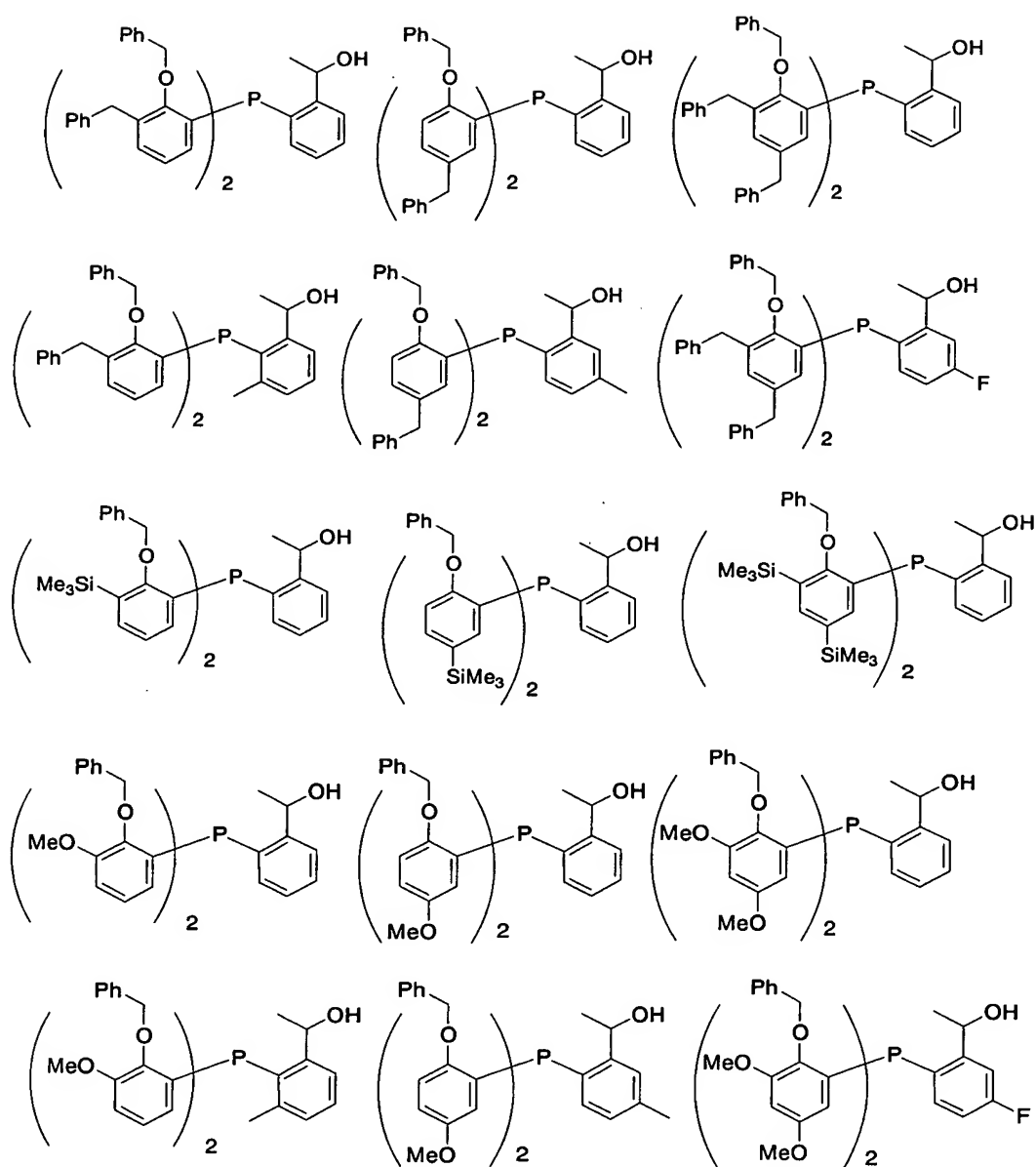


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The phosphine compound of formula (24B) is produced by reacting the phosphine compound of formula (24C) with a metal hydride compound or the organic metal compound of formula (24D),
 10 $R^{16}-Y$, wherein R^{16} and Y are as described above.

The molar ratio between the phosphine compound of formula (24C) and the metal hydride compound or the organic metal compound or the organic metal compound of formula (24D) in the reaction is not

particularly restricted, and the ratio is preferably in the range of 1:0.1 to 1:10, more preferably in the range of 1:0.5 to 1:5.

Examples of the metal hydride compound in the reaction above include, for example, sodium borohydride, potassium
5 borohydride, zinc borohydride, sodium cyanoborohydride, sodium triethylborohydride, lithium aluminum hydride, diisobutyl aluminum hydride, tri(tert-butoxy)lithium aluminum hydride and the like.

Examples of the organic metal compound of formula (24D)
10 include, for example, organic alkali metal compounds including organic lithium compounds such as methyl lithium, ethyl lithium, n-butyl lithium, sec-butyl lithium, tert-butyl lithium, lithium trimethylsilyl acetylide, lithium acetylide, trimethylsilylmethyl lithium, vinyl lithium, phenyl lithium and
15 allyl lithium; and organic alkaline earth metal halide such as organic magnesium halide including methylbromo magnesium, ethylbromo magnesium, phenylbromo magnesium, tolylbromo magnesium, benzylbromo magnesium or the like.

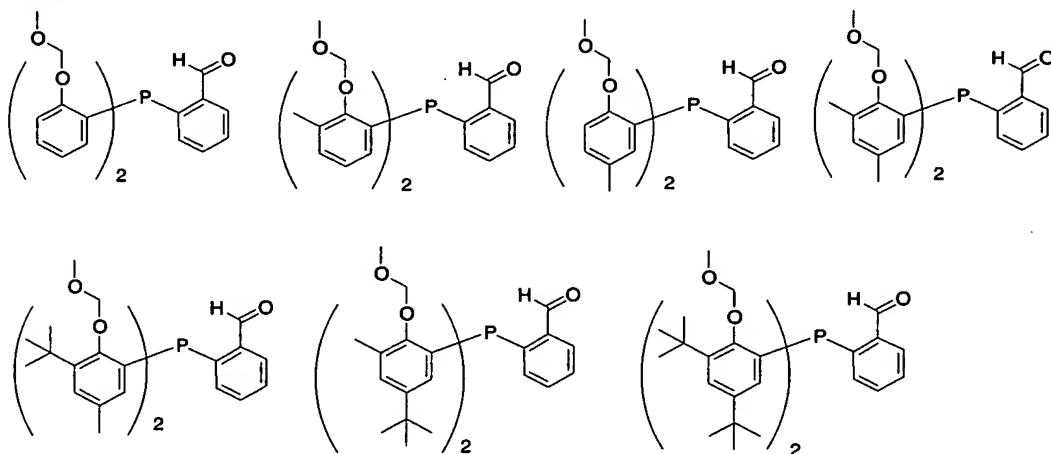
The reaction above is usually performed in an organic
20 solvent. Examples of the solvent include, for example, aprotic solvents including aromatic hydrocarbon solvents such as benzene toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; ether solvents such as diethyl ether, tetrahydrofuran, 1,4-dioxane or the like; amide solvents such
25 as hexamethyl phosphoric amide, dimethylformamide or the like; polar solvents such as acetonitrile, propionitrile, acetone, diethyl ketone, methyl isobutyl ketone, cyclohexanone or the like; and halogenated solvents such as dichloromethane, dichloroethane, chlorobenzene, dichlorobenzene or the like; and

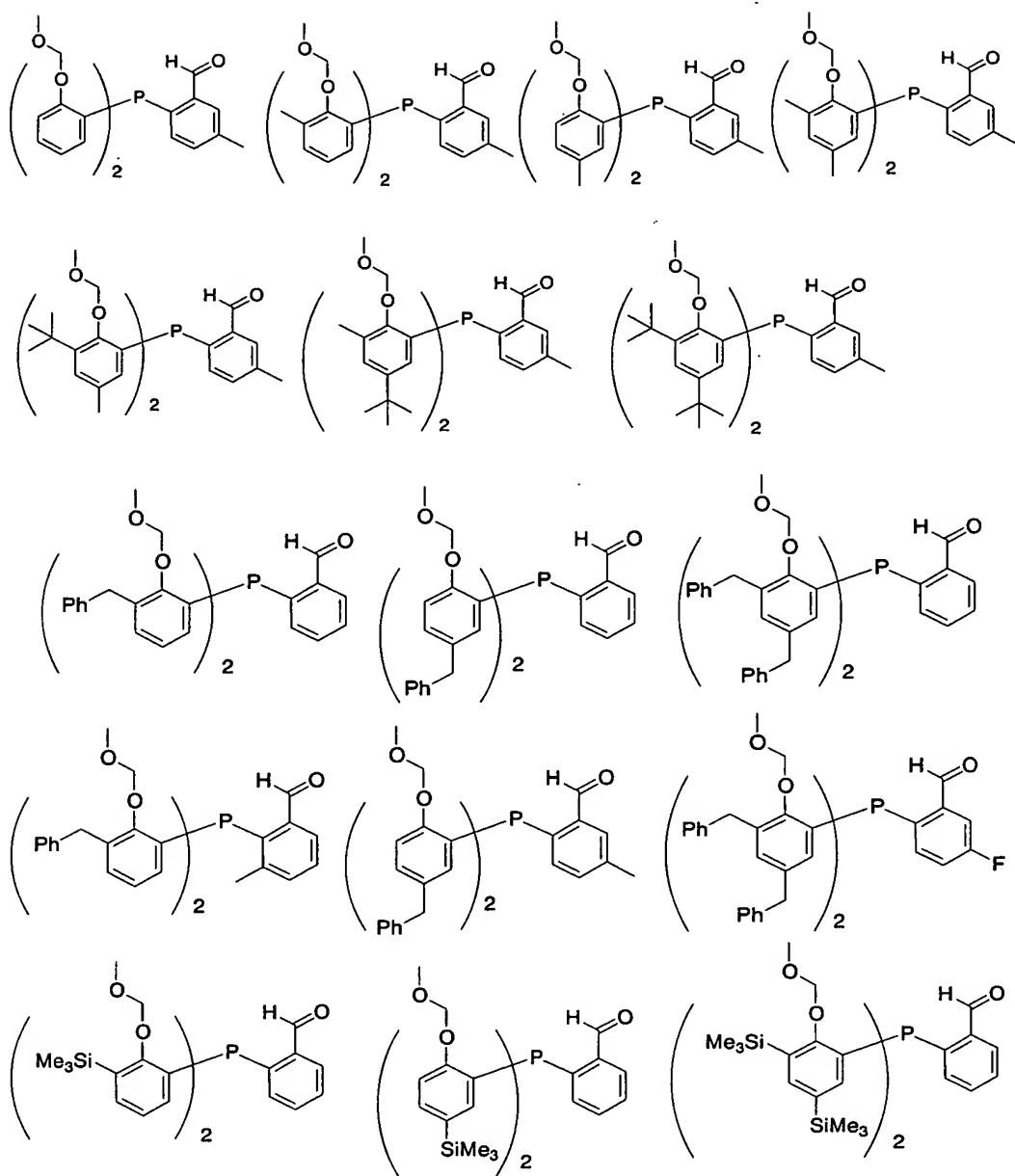
protonic solvents such as methanol, ethanol, isopropanol, butanol or the like. These solvents may be used alone or as a mixture of at least two of them. The amount thereof is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the phosphine compound of formula (24C).

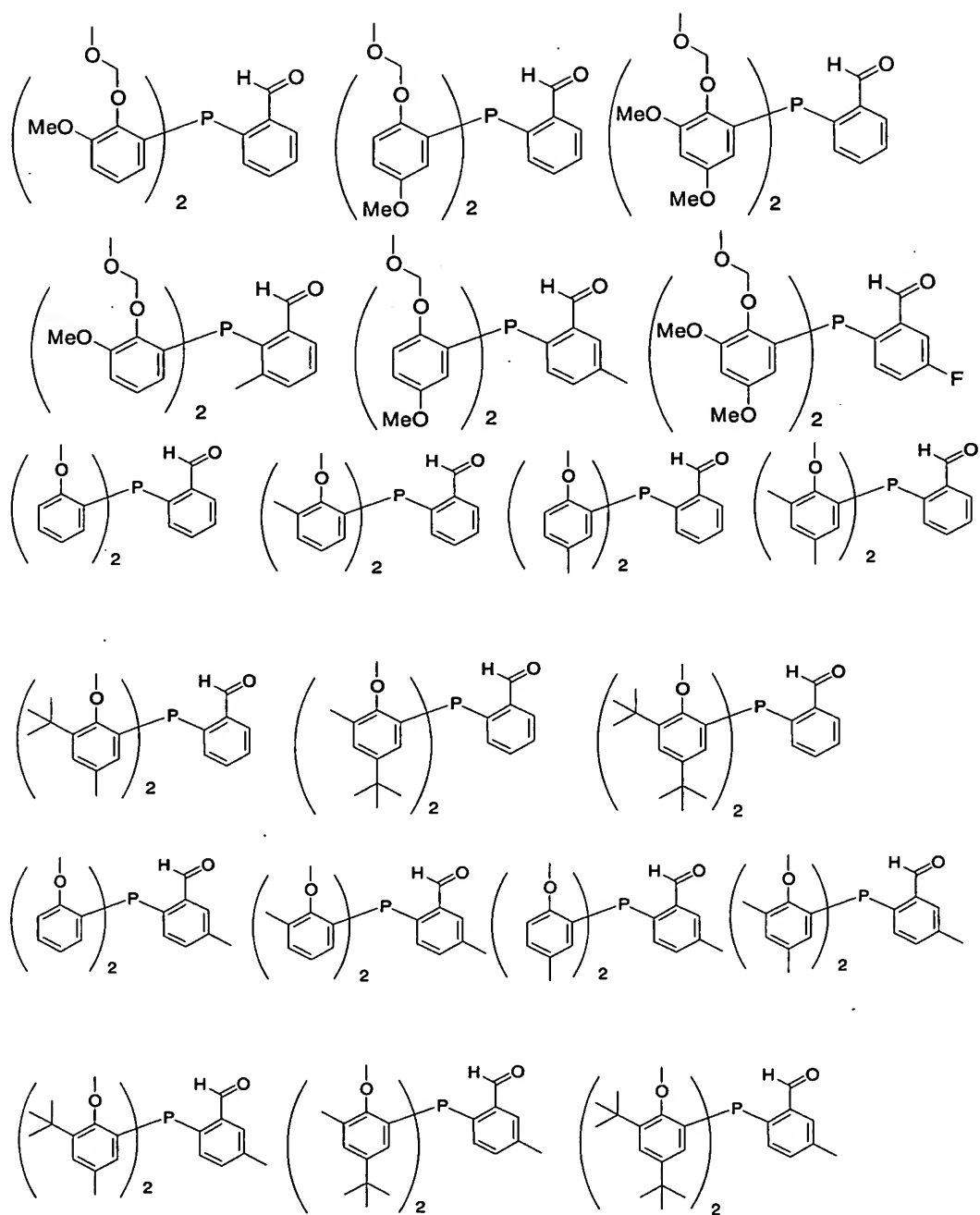
The reaction temperature is usually in the range of from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C.

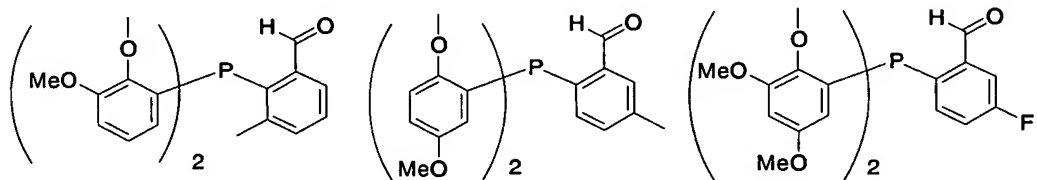
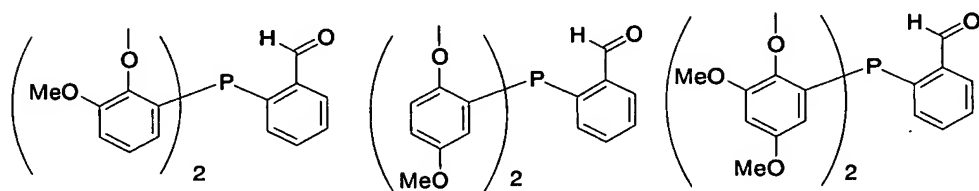
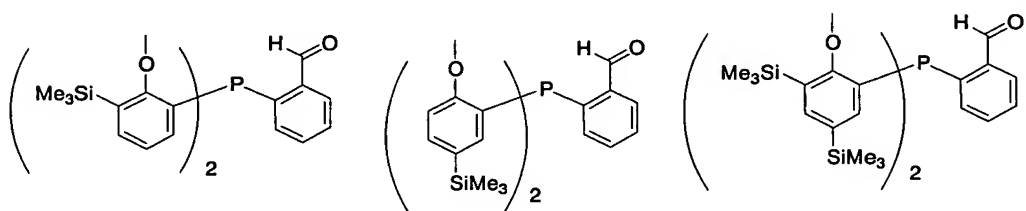
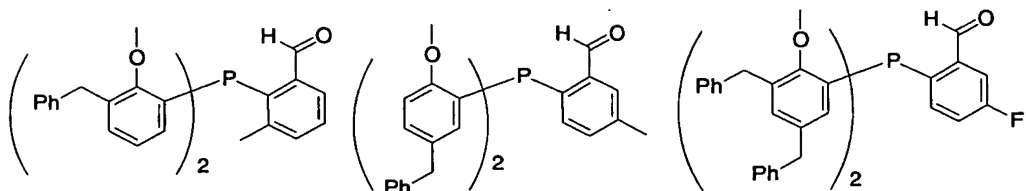
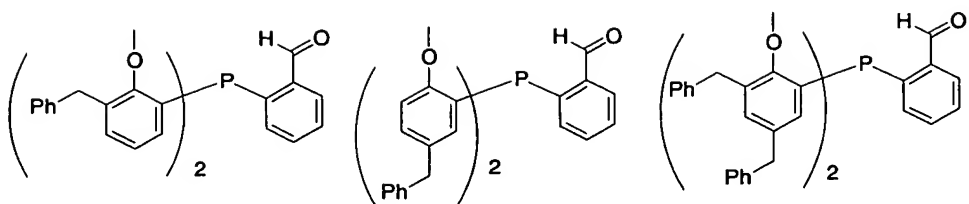
After the reaction, the phosphine compound of formula (24B) can be obtained from the reaction mixture by a conventional method, such as removing the solvent by evaporation. The reaction product can be purified by silica gel chromatography, if necessary.

Examples of the compound of formula (24C) include, for example, the following compounds:

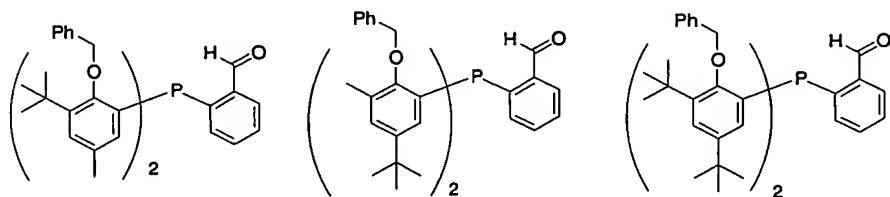
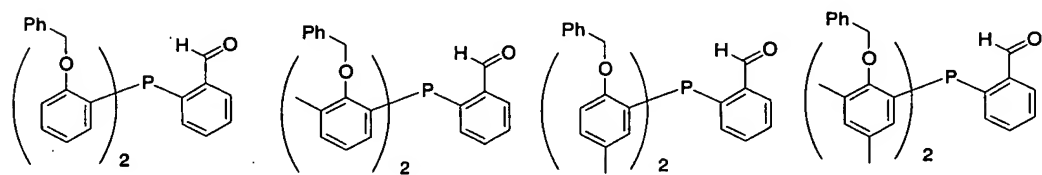


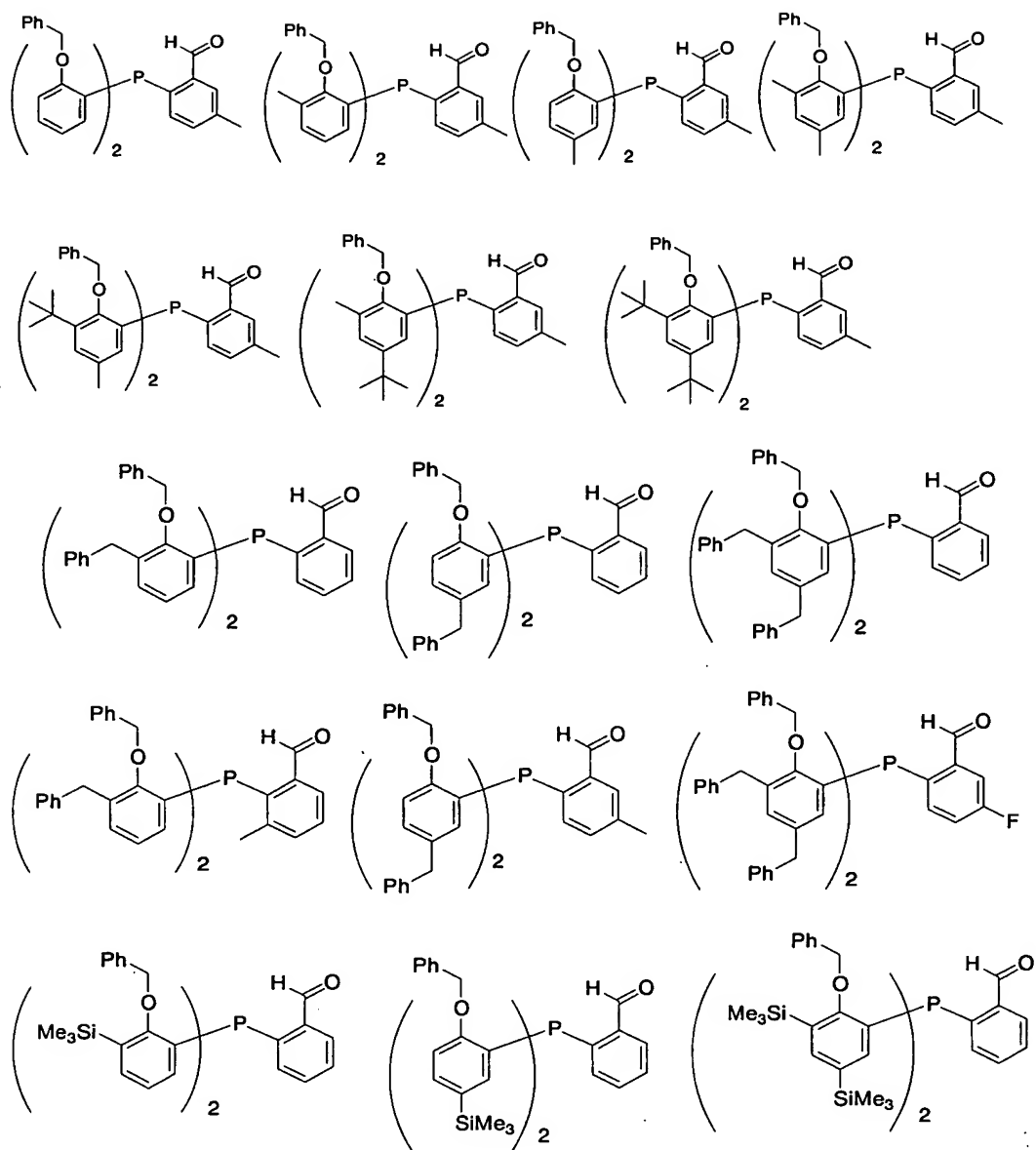


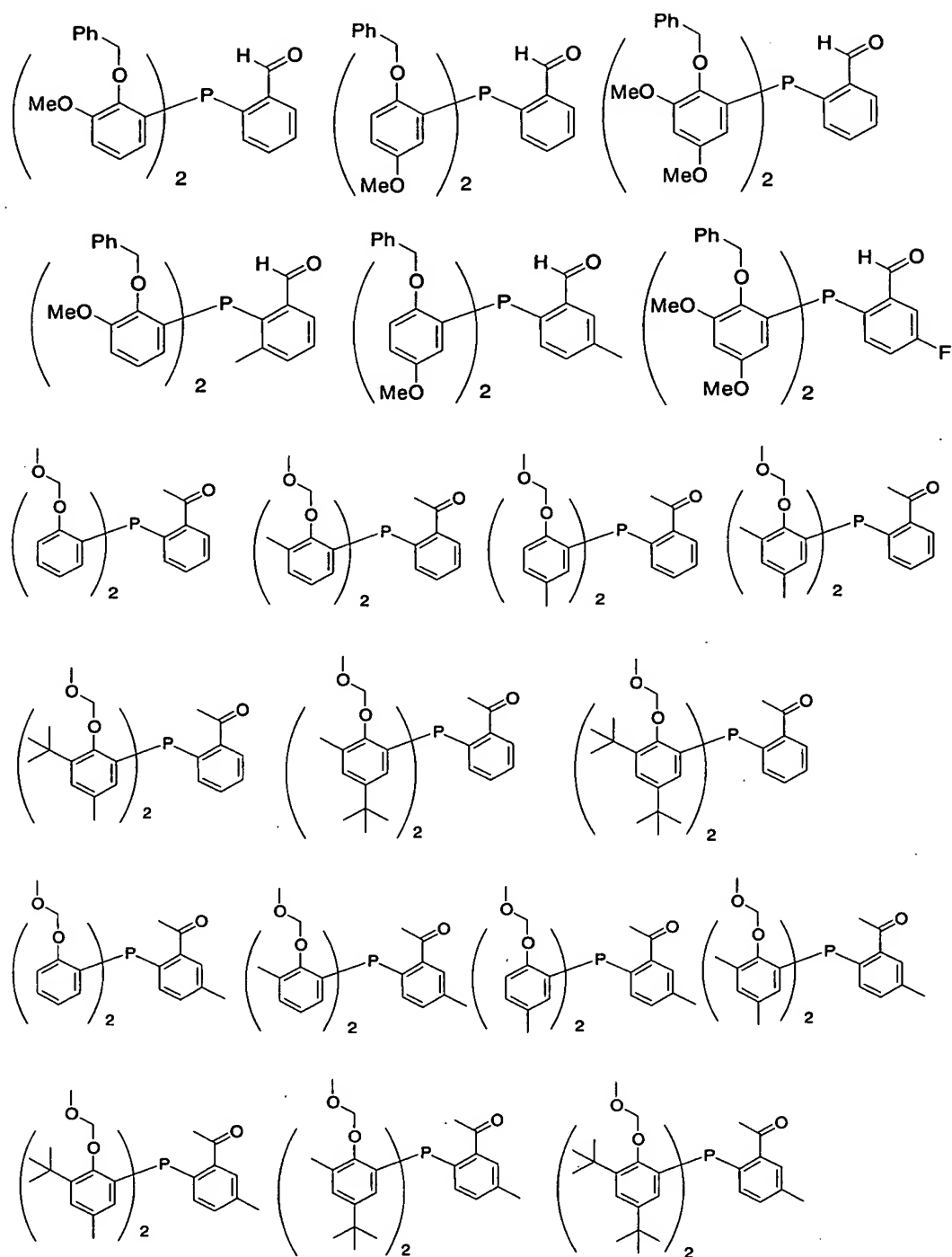




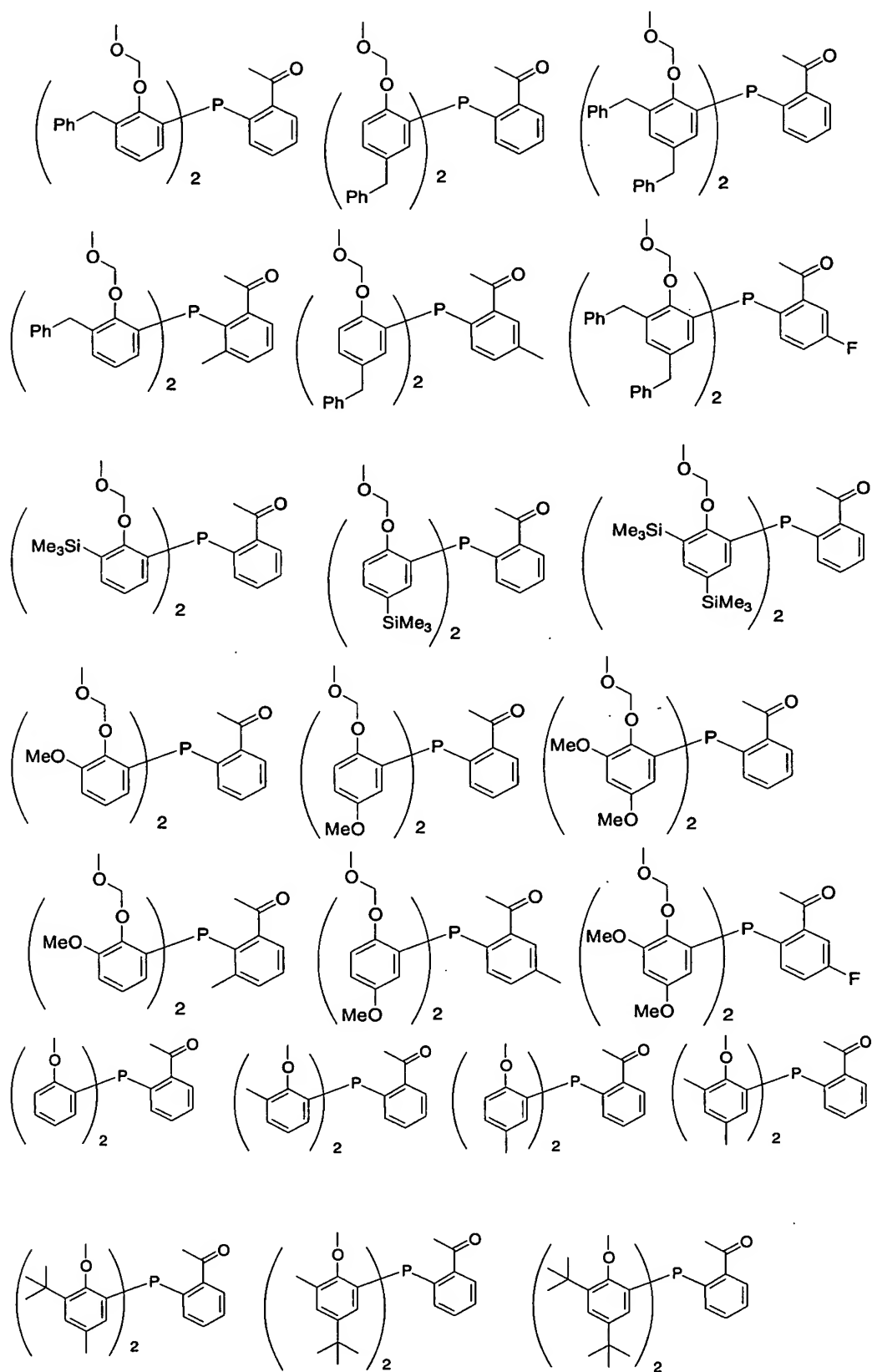
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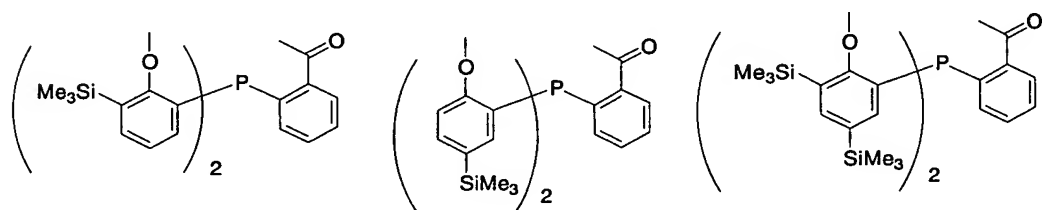
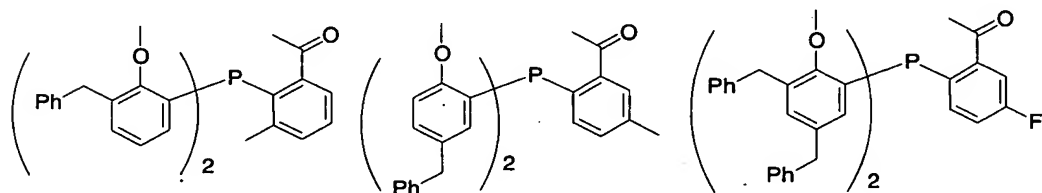
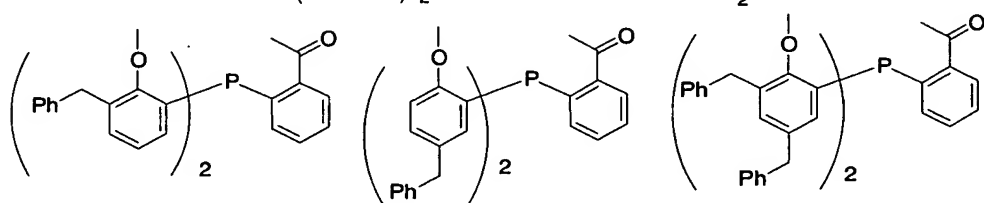
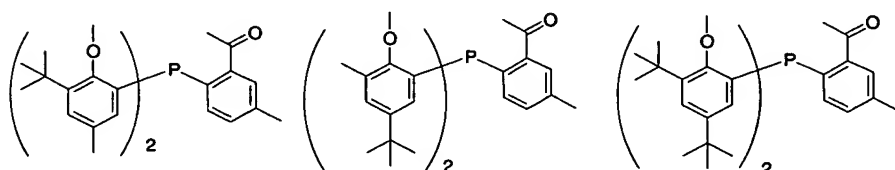
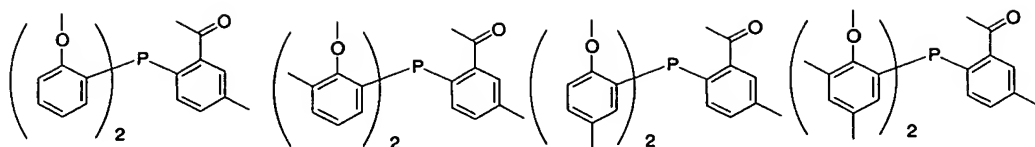




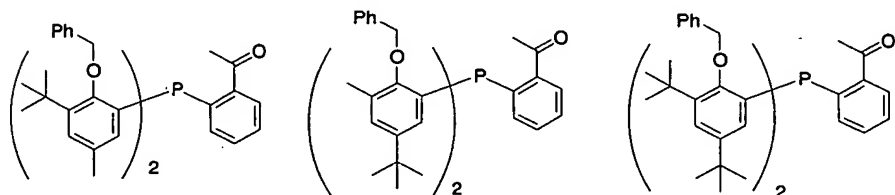
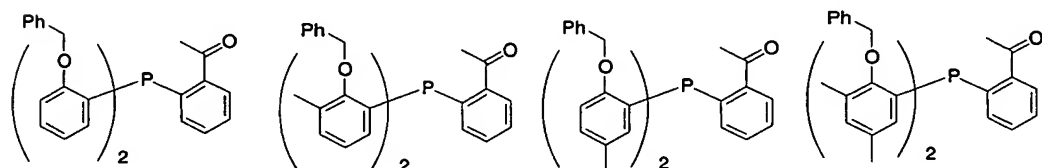


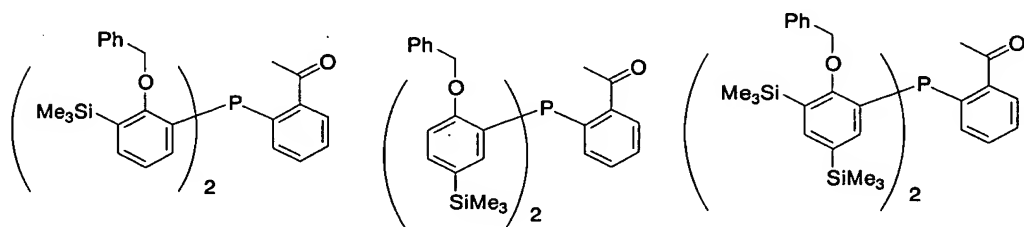
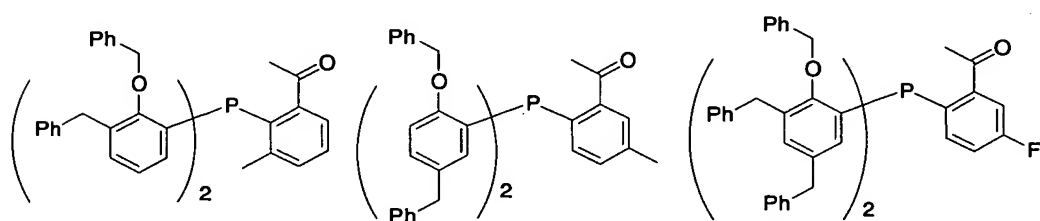
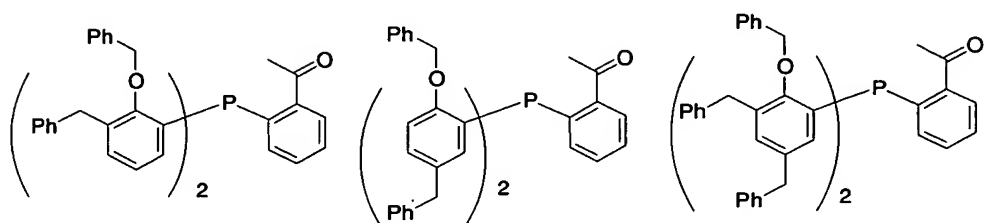
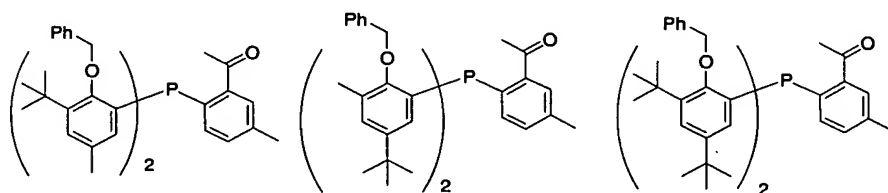
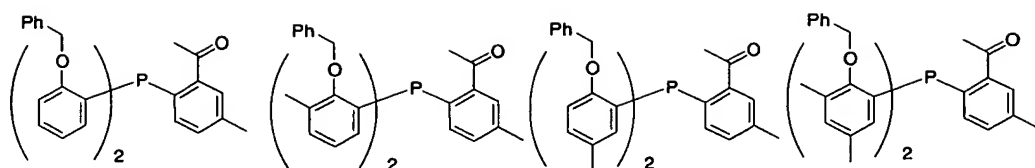
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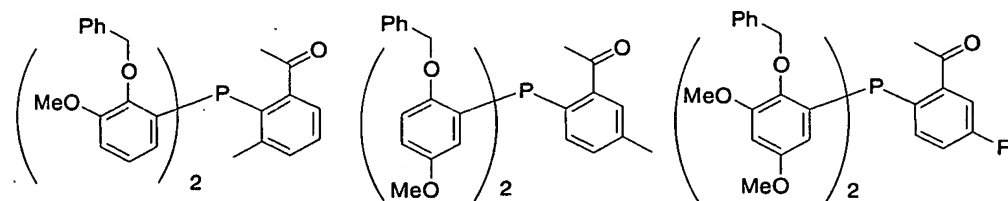
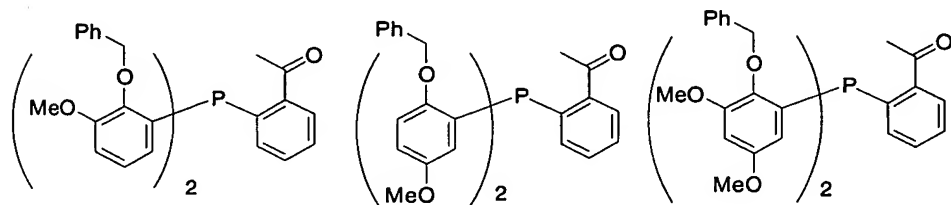


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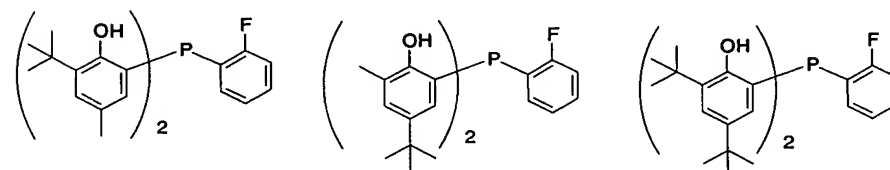
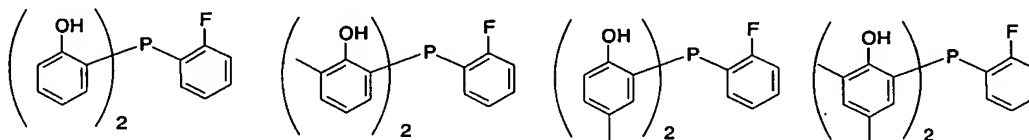




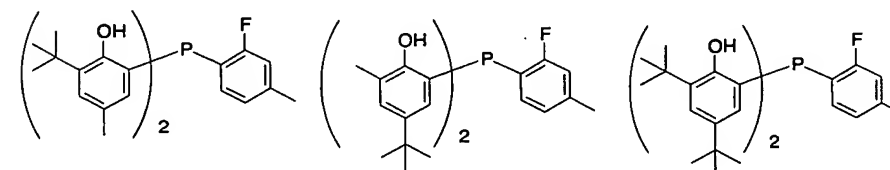
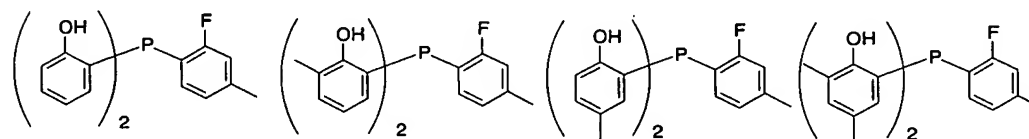
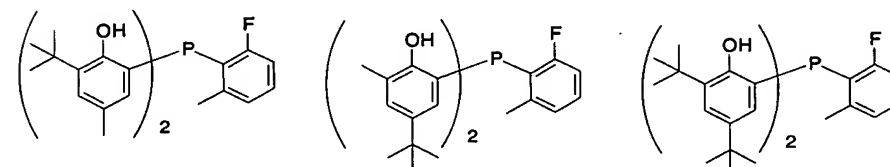
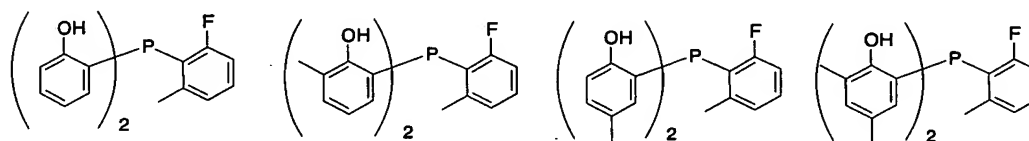
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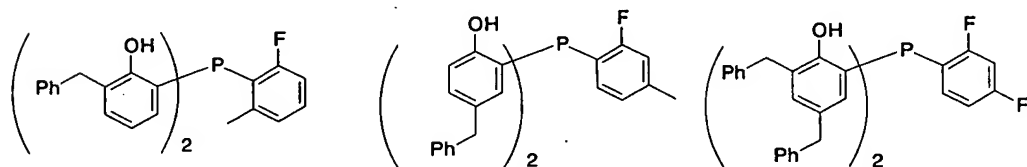
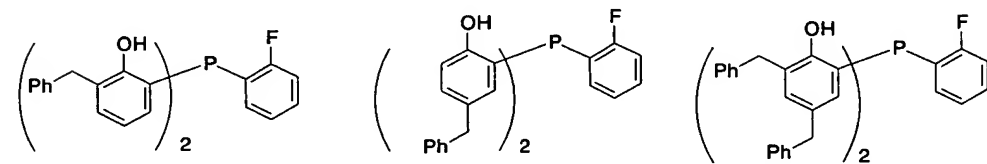
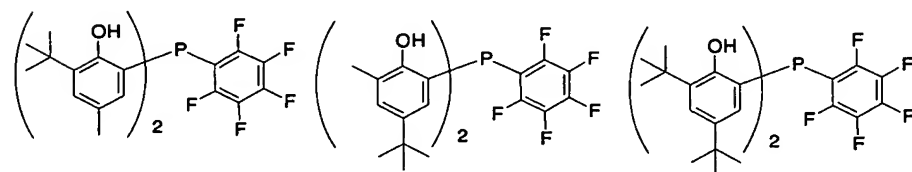
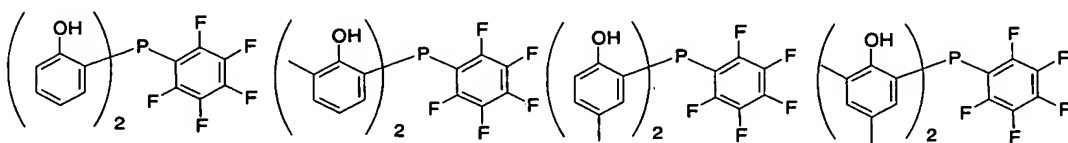
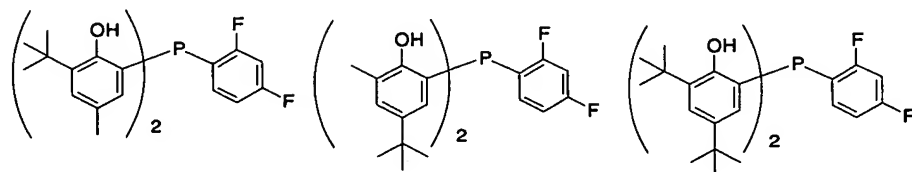
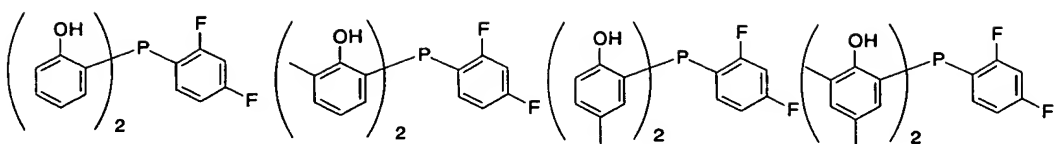


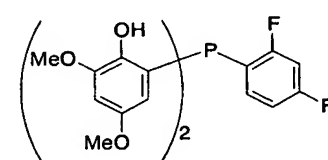
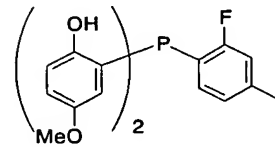
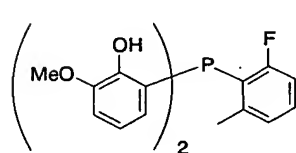
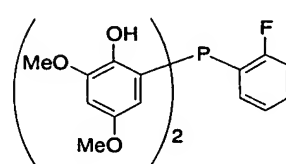
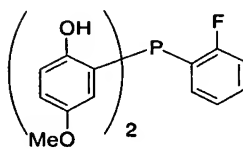
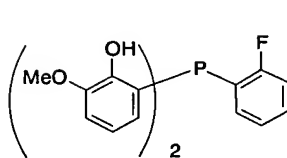
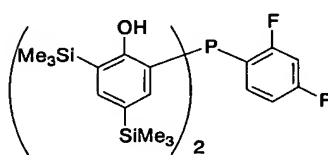
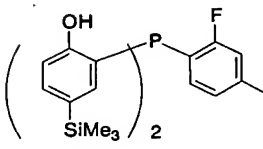
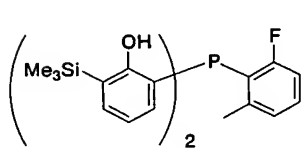
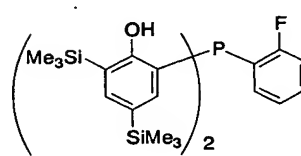
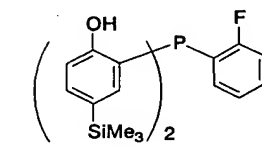
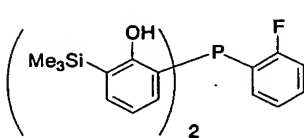
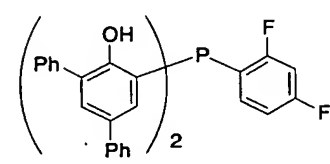
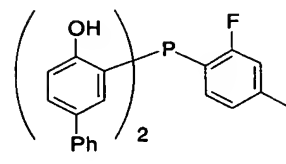
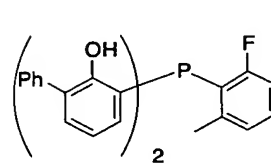
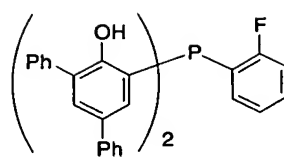
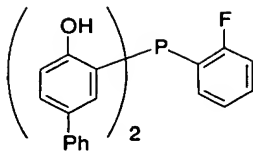
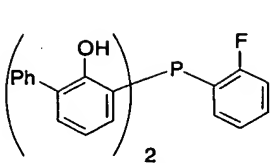
Examples of the compound of formula (1) wherein G^2 in is G^{25} , which corresponds to the compound of formula (25A) include, for example, the following compounds:

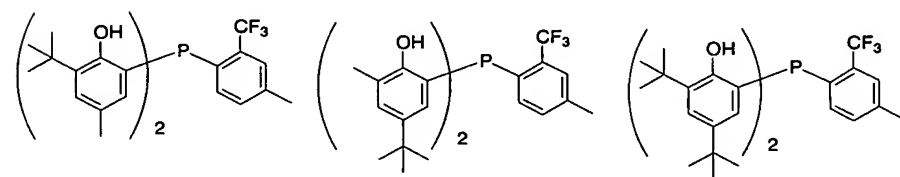
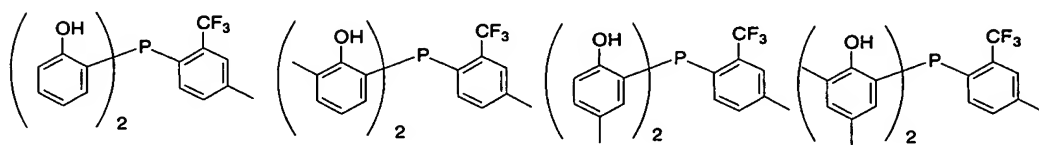
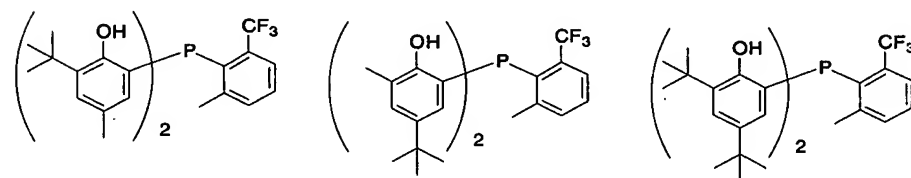
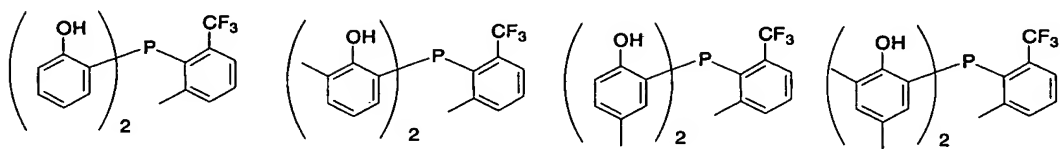
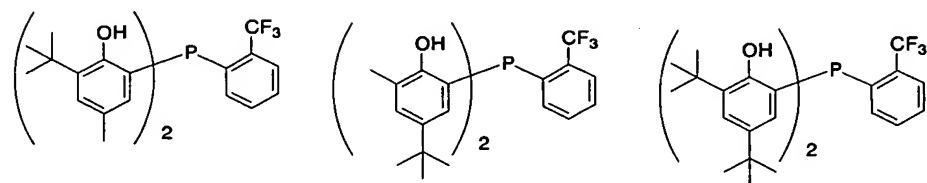
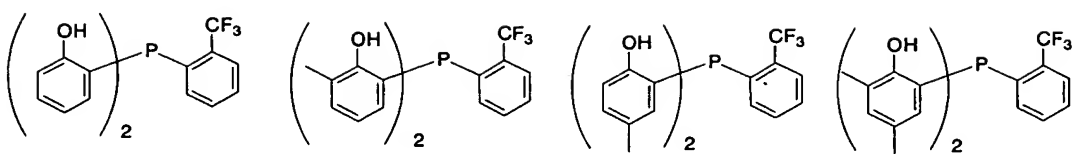


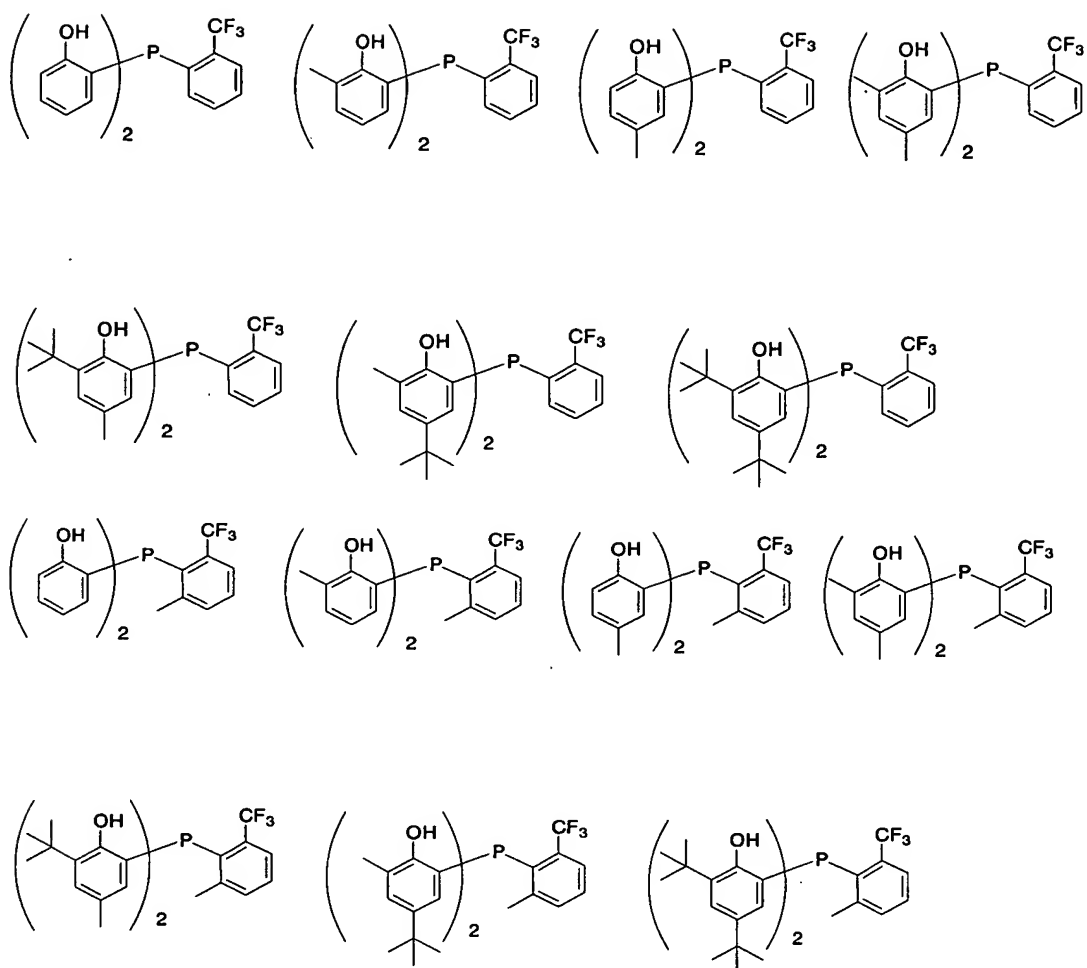
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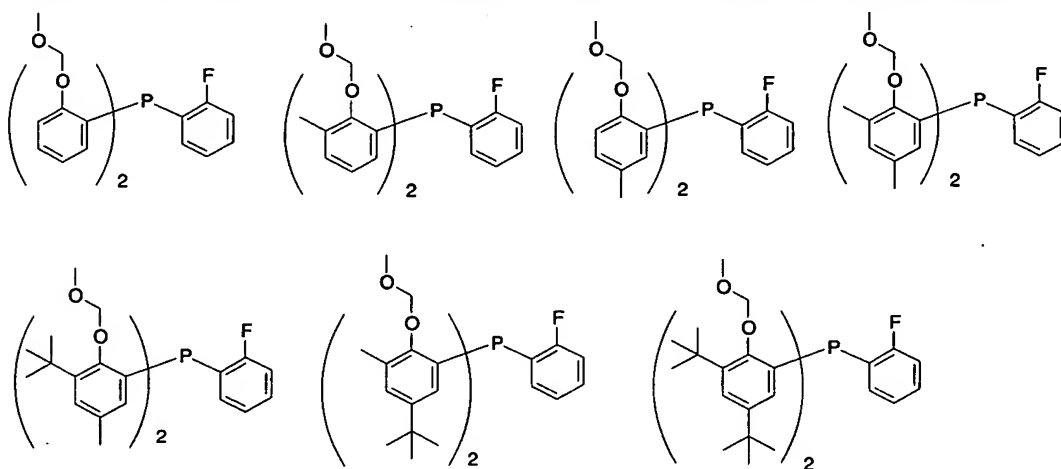


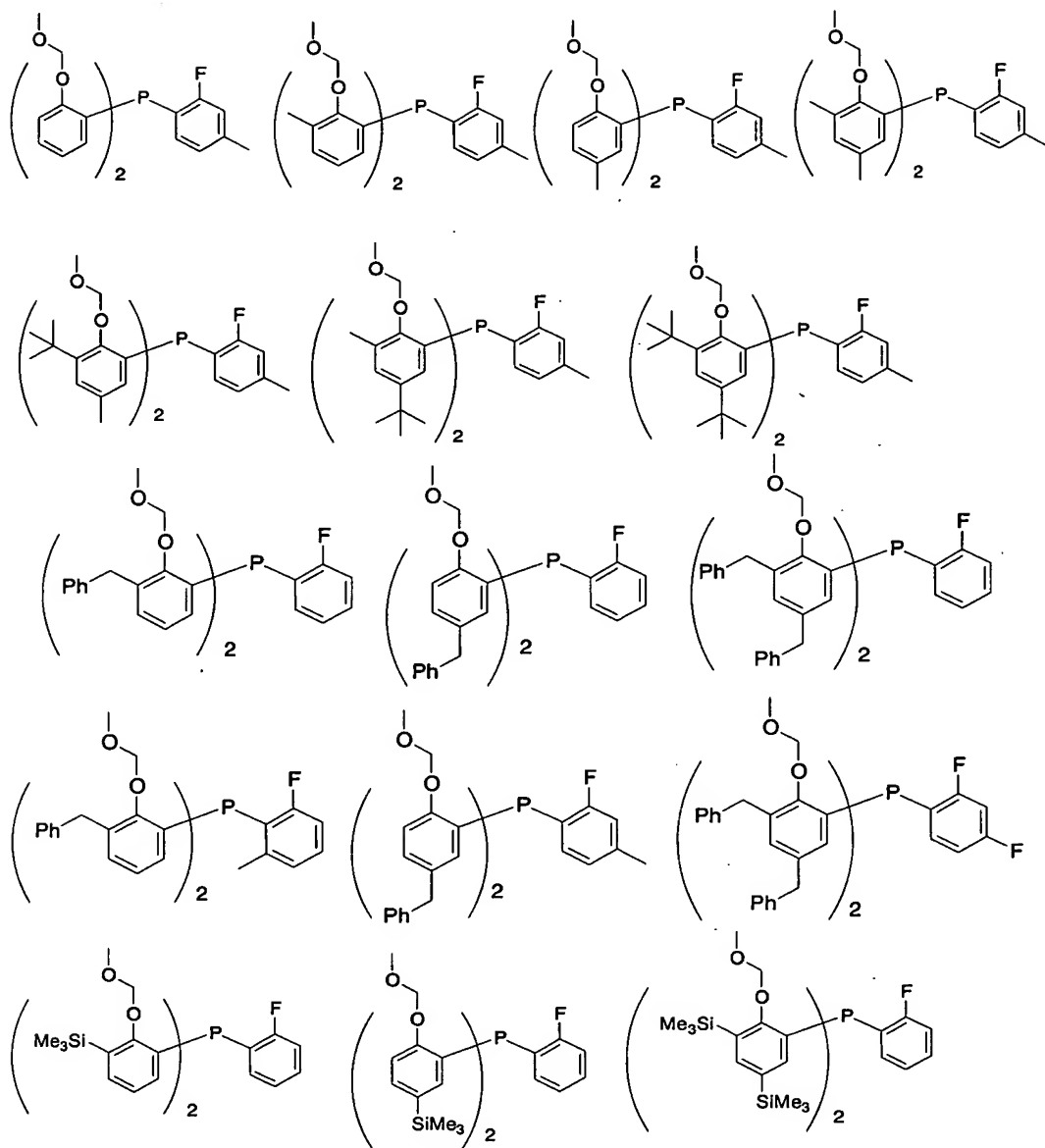


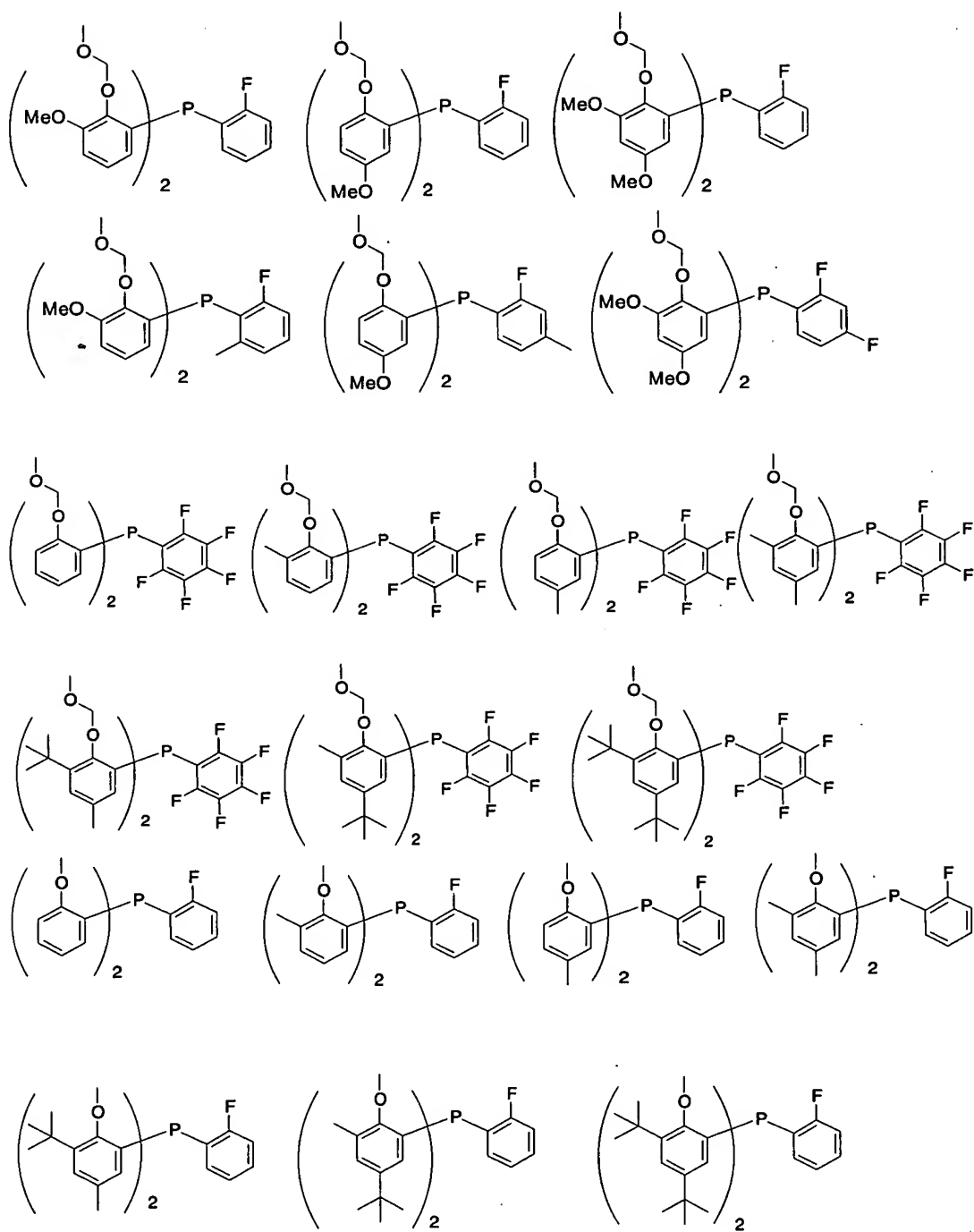


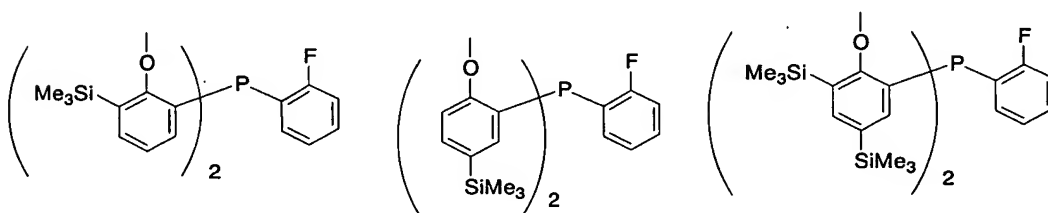
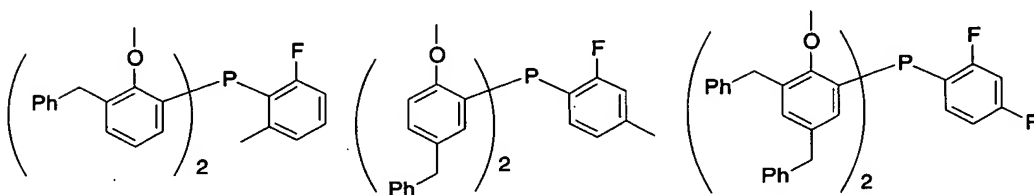
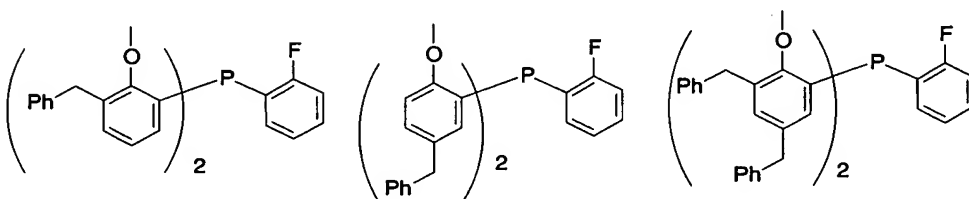
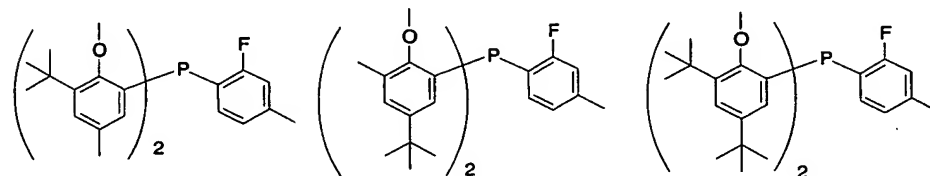
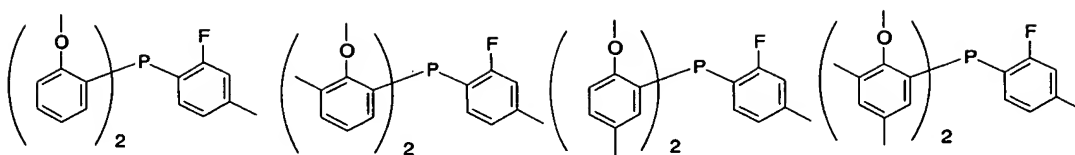


Examples of the compound of formula (1) wherein G^2 represents G^{25} , which corresponds to the compound of formula 5 (25B) include, for example, the following compounds:

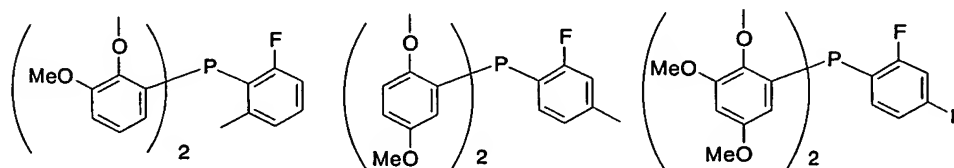
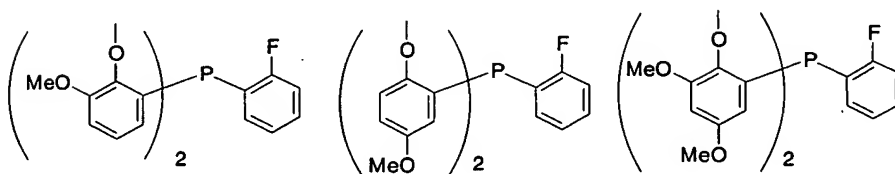


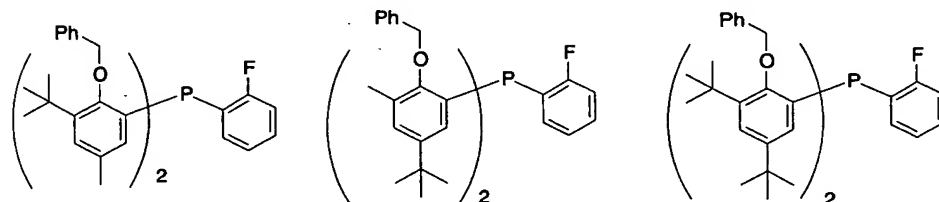
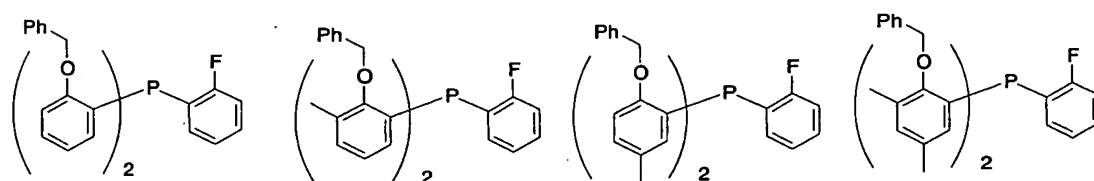
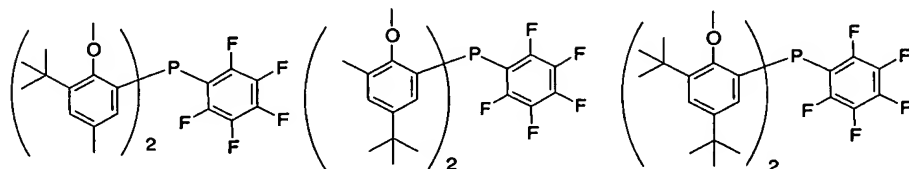
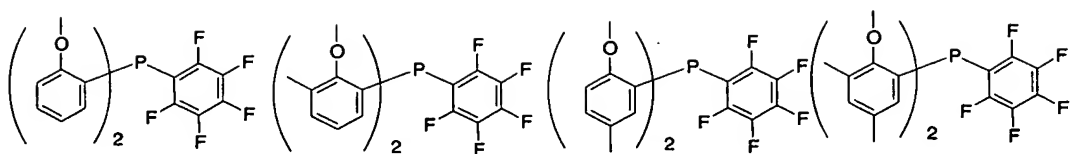




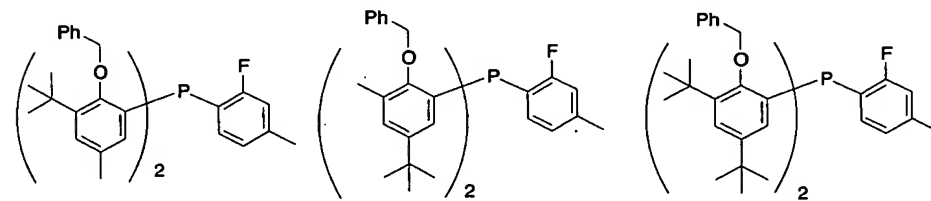
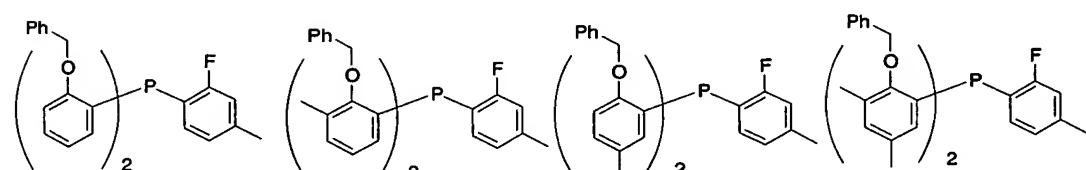


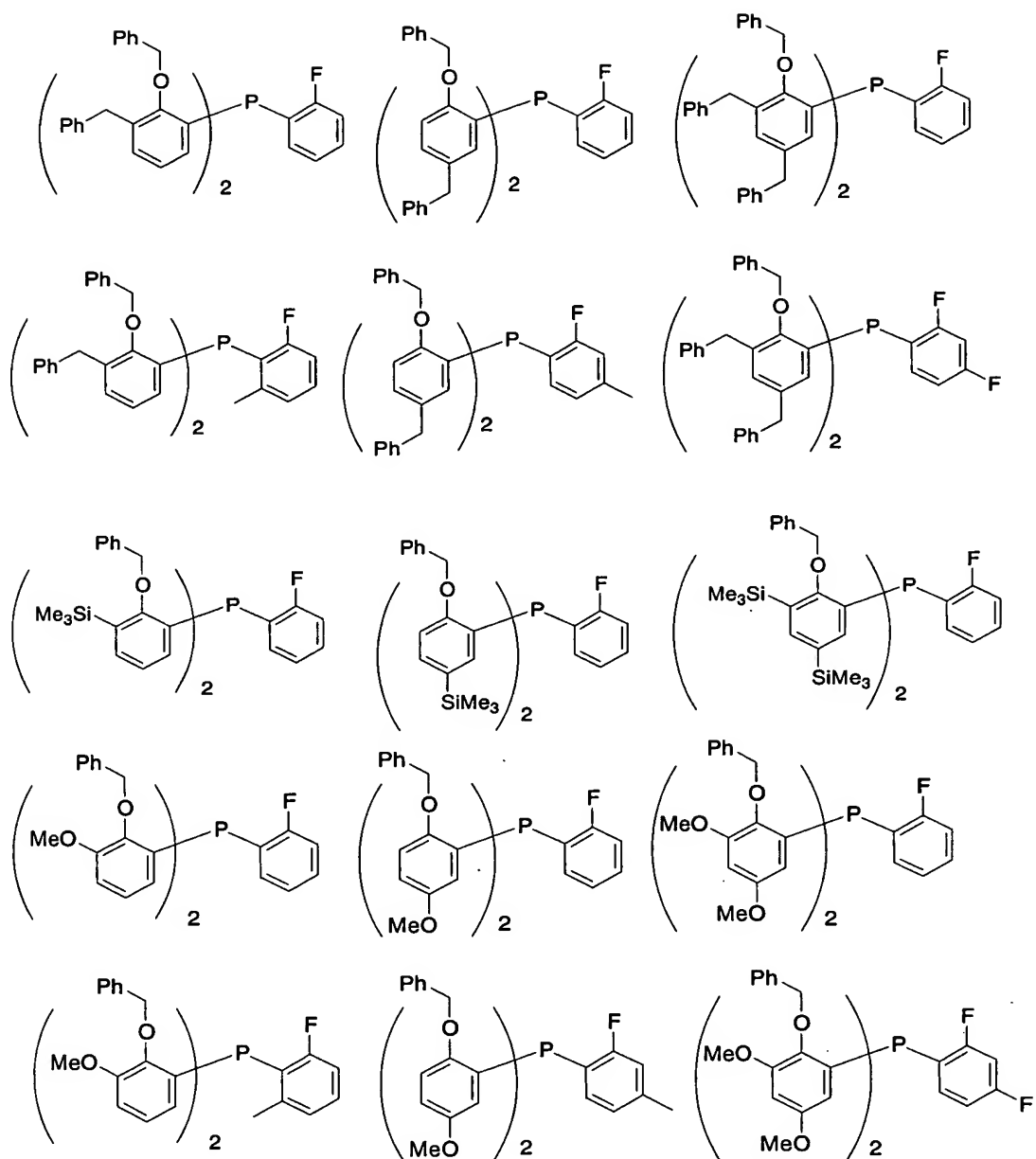
5

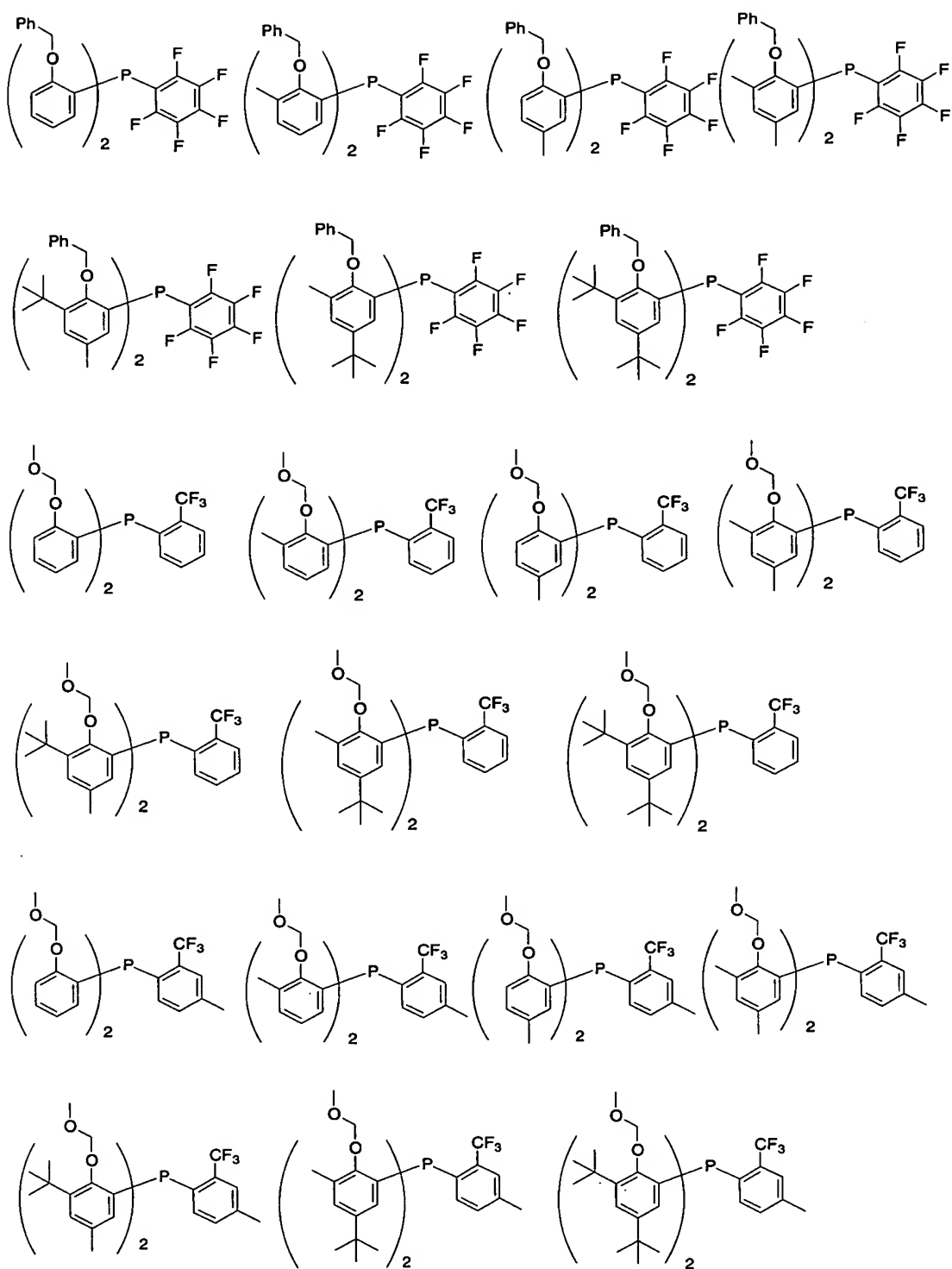




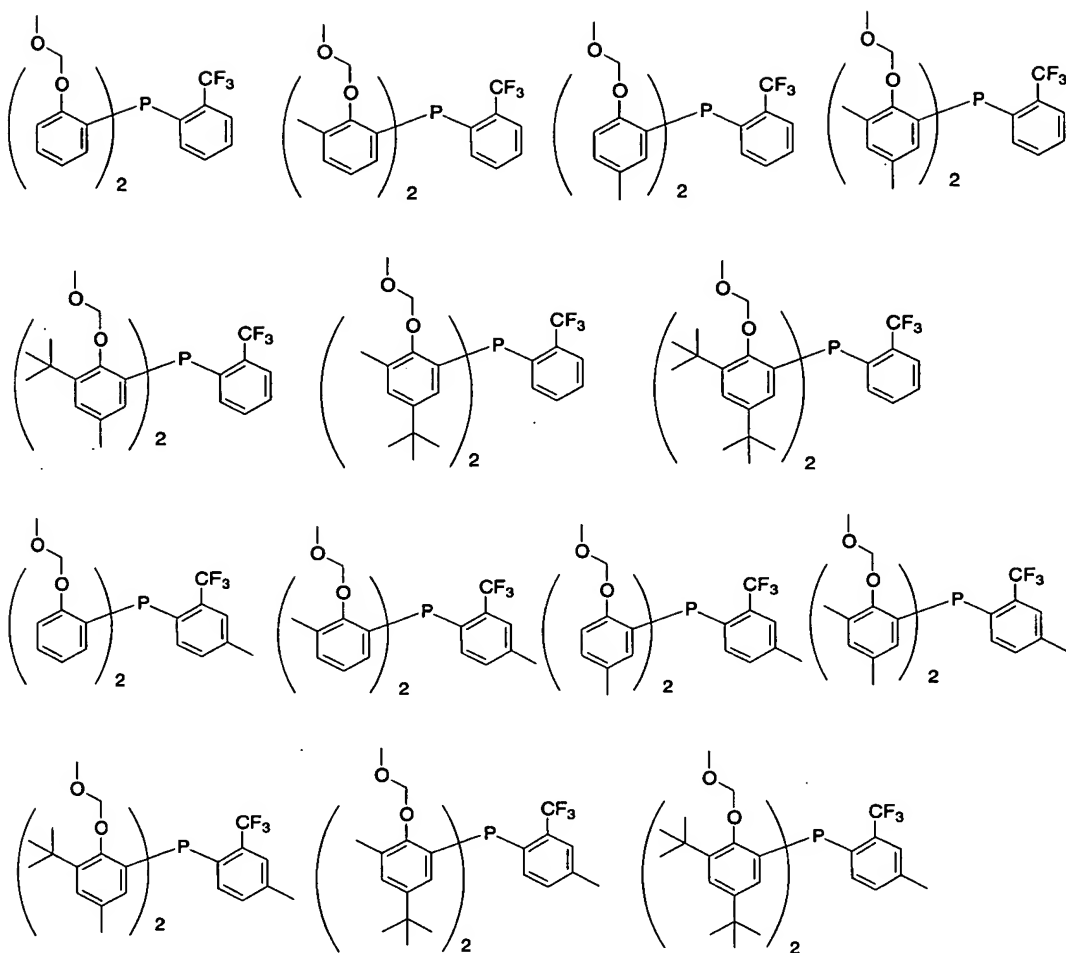
5







5



5 The phosphine compound of formula (25B) can be produced by reacting the phosphine halide compound of formula (25C) with the metal aryl compound of formula (25D). The molar ratio between the phosphine halide compound of formula (25C) and the metal aryl compound of formula (25D) is not particularly restricted,
 10 and the ratio is preferably in the range of 1:0.1 to 1:10, more preferably 1:0.5 to 1:5.

 Specific examples of the halogen atom represented by X² in the compound of formula (25C) or (25D) include fluorine, chlorine, bromine and iodine atoms. Chlorine atom is preferred.

15 Specific examples of the alkali metal represented by D

in the metal aryl compound include lithium, sodium and potassium atoms, and lithium atom is preferable among them. Specific examples of the alkaline earth metal represented by J include magnesium and calcium atoms, and magnesium atom is preferable.

5 The reaction above is usually performed in a solvent inert to the reaction. Examples of the solvent include, for example, aprotic solvents including aromatic hydrocarbon solvents such as benzene and toluene; aliphatic hydrocarbon solvents such as hexane, heptane or the like; ether solvents such as diethyl ether,
10 tetrahydrofuran, 1,4-dioxane or the like; polar solvents such as acetonitrile, acetone, diethyl ketone, methyl isobutyl ketone, cyclohexanone, ethyl acetate or the like; and halogenated solvents such as dichloromethane, dichloroethane, chlorobenzene, dichlorobenzene or the like; and protonic
15 solvents such as methanol, ethanol, isopropanol, butanol or the like. These solvents may be used alone or as a mixture of at least two of them. The amount thereof is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (6).

20 The reaction temperature is usually in the range of from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C.

 The phosphine compound of formula (3) can be obtained from the reaction mixture by a conventional method such as removing
25 the solvent by evaporation. The reaction product can be purified by recrystallization and silica gel chromatography, if necessary.

 The phosphine compound of formula (25B) can be produced by reacting phosphine dihalide of formula (25E) with the metal

aryl compound of formula (25F).

The molar ratio between phosphine dihalide of formula (25E) and the metal aryl compound of formula (25F) is not particularly limited, and it preferably in the range of 1:0.1 to 1:10, more preferably in the range of 1:1.5 to 1:5.

Specific examples of the halogen atom represented by X^2 in the compound of formula (25E) or (25F) include fluorine, chlorine, bromine and iodine atoms, and chlorine atom is preferable.

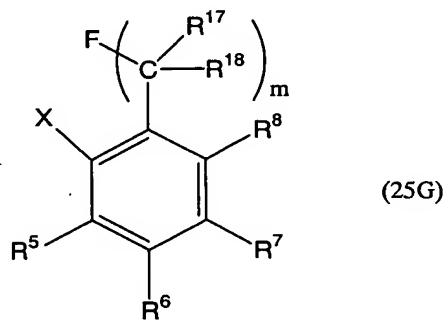
The reaction above is usually performed in a solvent inert to the reaction. Examples of the solvent include aprotic solvents including aromatic hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; ether solvents such as diethyl ether, tetrahydrofuran, 1,4-dioxane or the like; polar solvents such as acetonitrile, propionitrile, acetone, diethyl ketone, methyl isobutyl ketone, cyclohexanone, ethyl acetate or the like; and halogenated solvents such as dichloromethane, dichloroethane, chlorobenzene, dichlorobenzene or the like; and protonic solvents such as methanol, ethanol, isopropyl alcohol, butanol or the like. These solvents may be used alone or as a mixture of at least two of them. The amount thereof is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (25F).

The reaction temperature is usually in the range of from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C .

The phosphine compound of formula (25E) can be obtained from the reaction mixture by a conventional method such as

removing the solvent by evaporation. The reaction product can be purified by silica gel chromatography, if necessary.

The metal aryl compound of formula (25D) can be produced by reacting a fluorine-containing compound of formula (25G):



wherein R^5 , R^6 , R^7 , R^8 , R^{17} and R^{18} , and X and m are as described above, with, for example, a lithiating agent or magnesium metal.

The molar ratio between the metal aryl compound of formula (25G) and the lithiating agent or magnesium metal is not particularly restricted, and it is preferably in the range of 1:0.1 to 1:5, more preferably in the range of 1:0.5 to 1:2.5.

Examples of the lithiating agent include methyl lithium, n-butyl lithium, s-butyl lithium, t-butyl lithium, phenyl lithium and the like, and n-butyl lithium is preferable.

15 Specific examples of the halogen represented by X in the fluorine-containing compound of formula (25G) include fluorine, chlorine, bromine and iodine atoms, preferably chlorine atom.

The reaction above is usually performed in a solvent inert to the reaction. Examples of the solvent include aromatic hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture of at least two of them. The amount thereof is usually 1 to

200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the fluorine-containing compound of formula (25G).

5 The reaction can be performed by adding, for example, the lithiating agent or magnesium metal to the fluorine-containing compound of formula (25G). The reaction temperature is usually in the range of from -100°C or more to the boiling point or less of the solvent, and preferably in the range of -100°C to 100°C .

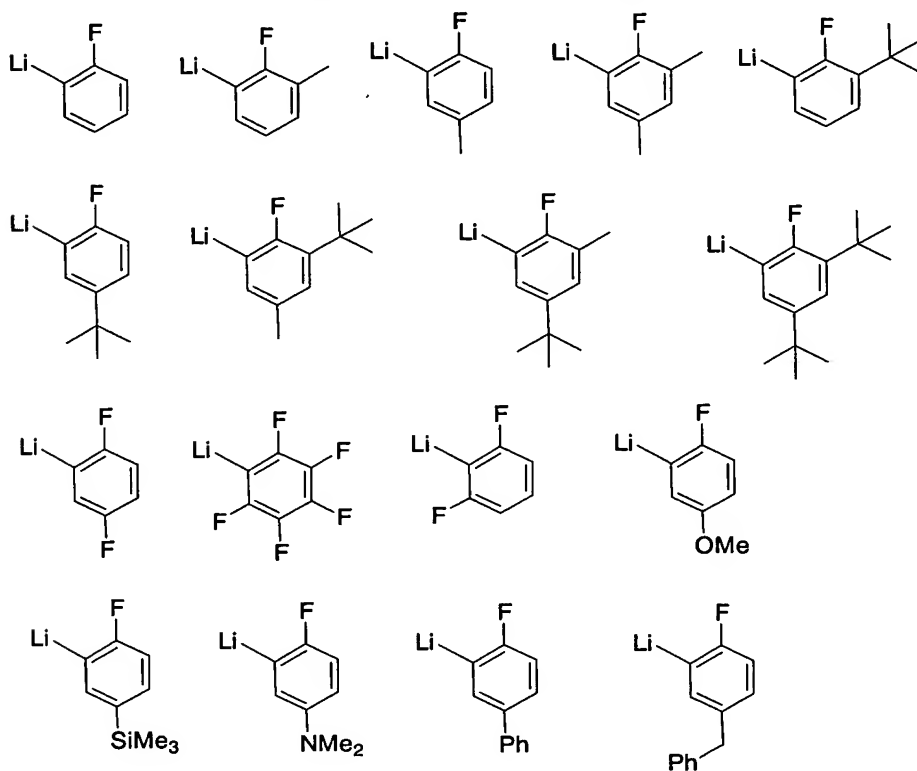
10 The phosphine dihalide of formula (25E) can be produced by reacting the phosphine dihalide represented by $\text{P}(\text{X}^2)_3$, wherein X^2 represents a halogen atom, with the metal aryl compound of formula (25D). The molar ratio between the phosphine halide and the metal aryl compound of formula (25D) is not particularly restricted, and the ratio is preferably in the range of 1:0.1
15 to 1:5, more preferably in the range of 1:0.5 to 1:2.

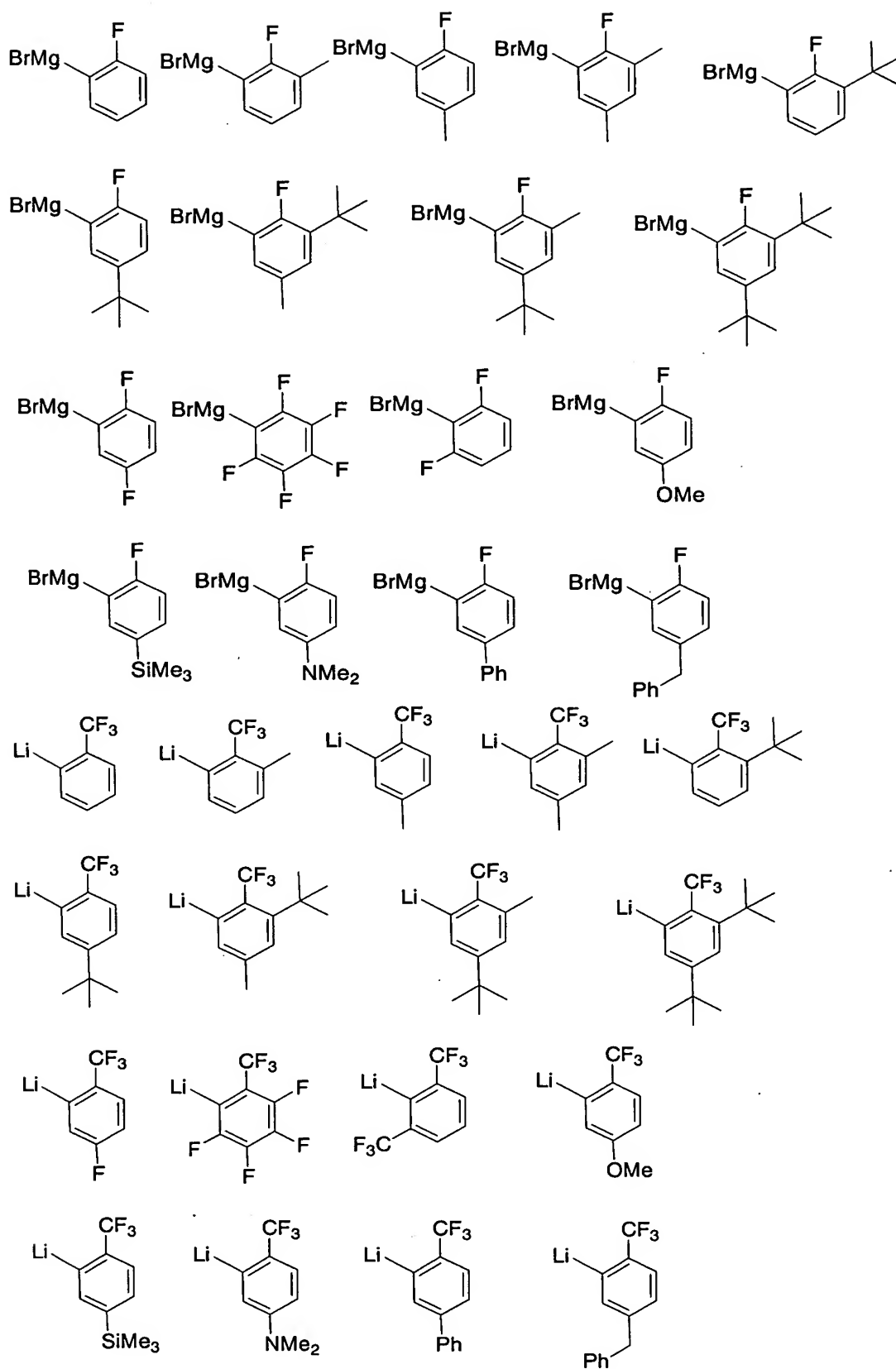
The reaction above is usually performed in a solvent inert to the reaction. Examples of the solvent include aromatic hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the
20 like; and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture at least two of them. The amount thereof is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (25D).

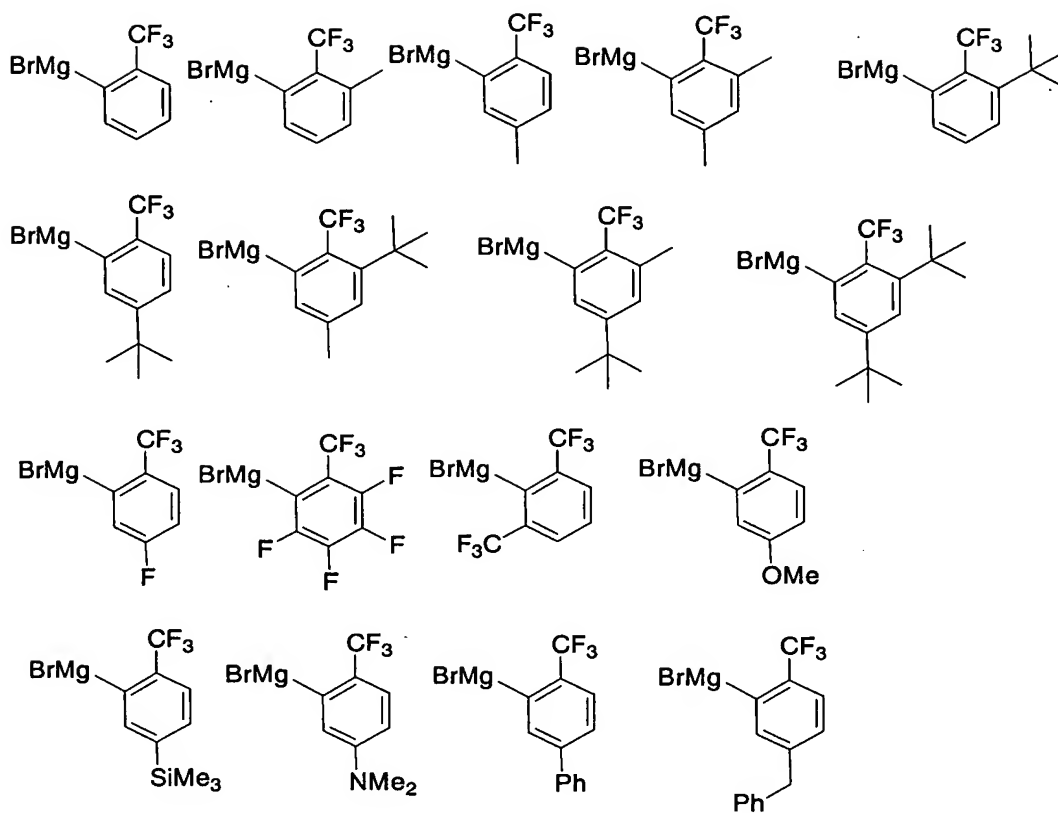
25 The reaction can be performed by adding, for example, phosphine trihalide to the metal aryl compound of formula (25D). The reaction temperature is usually in the range of from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C .

Phosphine dihalide of formula (25E) is obtained, for example, by removing insolubles by filtration followed by removing the solvent by evaporation. The product can be purified by, for example, distillation, if necessary.

5 Examples of the metal aryl compound of formula (25F) include, for example, the following compounds:







Examples of the phosphine dihalide of formula (25E) include, for example, the following compounds:

- 5 2-fluorophenyl dichlorophosphine, 2,6-difluorophenyl dichlorophosphine, 2,4,6-trifluorophenyl dichlorophosphine, pentafluorophenyl dichlorophosphine, 2-fluoro-6-methyl dichlorophosphine, 2-fluoro-6-tert-butylphenyl dichlorophosphine, 2-fluoro-4,6-dimethylphenyl
- 10 dichlorophosphine, 2-fluoro-4,6-di(tert-butyl)phenyl dichlorophosphine, 2-fluoro-4-methyl-6-tert-butylphenyl dichlorophosphine, 2-fluoro-5-phenylphenyl dichlorophosphine, 2-fluoro-4-phenylphenyl dichlorophosphine, 2-fluoro-6-phenylphenyl dichlorophosphine,

2-fluoro-5-benzylphenyl dichlorophosphine,
 2-fluoro-4-benzylphenyl dichlorophosphine,
 2-fluoro-6-benzylphenyl dichlorophosphine,
 2-fluoro-5-phenoxyphenyl dichlorophosphine,
 5 2-fluoro-4-phenoxyphenyl dichlorophosphine,
 2-fluoro-6-phenoxyphenyl dichlorophosphine,
 2-fluoro-5-methoxyphenyl dichlorophosphine,
 2-fluoro-4-methoxyphenyl dichlorophosphine,
 2-fluoro-6-methoxyphenyl dichlorophosphine,
 10 2-fluoro-5-trimethylsilylphenyl dichlorophosphine,
 2-fluoro-4-trimethylsilylphenyl dichlorophosphine,
 2-fluoro-6-trimethylsilylphenyl dichlorophosphine,
 2-fluoro-5-dimethylaminophenyl dichlorophosphine,
 2-fluoro-4-dimethylaminophenyl dichlorophosphine,
 15 2-fluoro-6-dimethylaminophenyl dichlorophosphine,
 2-fluorophenyl dibromophosphine, 2,6-difluorophenyl
 dibromophosphine, 2,4,6-trifluorophenyl dibromophosphine,
 pentafluorophenyl dibromophosphine, 2-fluoro-6-methylphenyl
 dibromophosphine, 2-fluoro-6-tert-butylphenyl
 20 dibromophosphine, 2-fluoro-4,6-di(tert-butyl)phenyl
 dibromophosphine, 2-fluoro-4-methyl-6-tert-butylphenyl
 dibromophosphine, 2-fluoro-5-phenylphenyl dibromophosphine,
 2-fluoro-4-phenylphenyl dibromophosphine,
 2-fluoro-6-phenylphenyl dibromophosphine,
 25 2-fluoro-5-benzylphenyl dibromophosphine,
 2-fluoro-4-benzylphenyl dibromophosphine,
 2-fluoro-6-benzylphenyl dibromophosphine,
 2-fluoro-5-phenoxyphenyl dibromophosphine,
 2-fluoro-4-phenoxyphenyl dibromophosphine,

2-fluoro-6-phenoxyphenyl dibromophosphine,
 2-fluoro-5-methoxyphenyl dibromophosphine,
 2-fluoro-4-methoxyphenyl dibromophosphine,
 2-fluoro-6-methoxyphenyl dibromophosphine,
 5 2-fluoro-5-trimethylsilylphenyl dibromophosphine,
 2-fluoro-4-trimethylsilylphenyl dibromophosphine,
 2-fluoro-6-trimethylsilylphenyl dibromophosphine,
 2-fluoro-5-dimethylaminophenyl dibromophosphine,
 2-fluoro-4-dimethylaminophenyl dibromophosphine,
 10 2-fluoro-6-dimethylaminophenyl dibromophosphine,
 2-trifluoromethylphenyl dibromophosphine,
 2,6-bistrifluoromethylphenyl dichlorophosphine,
 2,4,6-tristrifluoromethylphenyl dichlorophosphine,
 2-trifluoromethyl-6-methylphenyl dichlorophosphine,
 15 2-trifluoromethyl-6-tert-butylphenyl dichlorophosphine,
 2-trifluoromethyl-4,6-dimethylphenyl dichlorophosphine,
 2-trifluoromethyl-4,6-di(tert-butyl)phenyl
 dichlorophosphine,
 2-trifluoromethyl-4-methyl-6-tert-butylphenyl
 20 dichlorophosphine, 2-trifluoromethyl-5-phenylphenyl
 dichlorophosphine, 2-trifluoromethyl-4-phenylphenyl
 dichlorophosphine, 2-trifluoromethyl-6-phenylphenyl
 dichlorophosphine, 2-trifluoromethyl-5-benzylphenyl
 dichlorophosphine, 2-trifluoromethyl-4-benzylphenyl
 25 dichlorophosphine, 2-trifluoromethyl-6-benzylphenyl
 dichlorophosphine, 2-trifluoromethyl-5-phenoxyphenyl
 dichlorophosphine 2-trifluoromethyl-4-phenoxyphenyl
 dichlorophosphine, 2-trifluoromethyl-6-phenoxyphenyl
 dichlorophosphine, 2-trifluoromethyl-5-methoxyphenyl

dichlorophosphine, 2-trifluoromethyl-4-methoxyphenyl
dichlorophosphine, 2-trifluoromethyl-6-methoxyphenyl
dichlorophosphine, 2-trifluoromethyl-5-trimethylsilylphenyl
dichlorophosphine, 2-trifluoromethyl-4-trimethylsilylphenyl
5 dichlorophosphine, 2-trifluoromethyl-6-trimethylsilylphenyl
dichlorophosphine, 2-trifluoromethyl-5-dimethylaminophenyl
dichlorophosphine, 2-trifluoromethyl-4-dimethylaminophenyl
dichlorophosphine, 2-trifluoromethyl-6-dimethylaminophenyl
dichlorophosphine,
10 2-trifluoromethylphenyl dibromophosphine,
2,6-bistrifluoromethylphenyl dibromophosphine,
2,4,6-tristrifluoromethylphenyl dibromophosphine,
2-trifluoromethyl-6-methylphenyl dibromophosphine,
2-trifluoromethyl-6-tert-butylphenyl dibromophosphine,
15 2-trifluoromethyl-4,6-dimethylphenyl dibromophosphine,
2-trifluoromethyl-4,6-di(tert-butyl)phenyl dibromophosphine,
2-trifluoromethyl-4-methyl-6-tert-butylphenyl
dibromophosphine, 2-trifluoromethyl-5-phenylphenyl
dibromophosphine, 2-trifluoromethyl-4-phenylphenyl
20 dibromophosphine, 2-trifluoromethyl-6-phenylphenyl
dibromophosphine, 2-trifluoromethyl-5-benzylphenyl
dibromophosphine, 2-trifluoromethyl-4-benzylphenyl
dibromophosphine, 2-trifluoromethyl-6-benzylphenyl
dibromophosphine, 2-trifluoromethyl-5-phenoxyphenyl
25 dibromophosphine, 2-trifluoromethyl-4-phenoxyphenyl
dibromophosphine, 2-trifluoromethyl-6-phenoxyphenyl
dibromophosphine, 2-trifluoromethyl-5-methoxyphenyl
dibromophosphine, 2-trifluoromethyl-4-methoxyphenyl
dibromophosphine, 2-trifluoromethyl-6-methoxyphenyl

dibromohosphine, 2-trifluoromethyl-5-trimethylsilylphenyl
dibromophosphine, 2-trifluoromethyl-4-trimethylsilylphenyl
dibromophosphine, 2-trifluoromethyl-6-trimethylsilylphenyl
dibromophosphine, 2-trifluoromethyl-5-dimethylaminophenyl
5 dibromophosphine, 2-trifluoromethyl-4-dimethylaminophenyl
dibromophosphine, and
2-trifluoromethyl-6-dimethylaminophenyl dibromophosphine.

Examples of the fluorine-containing compound of formula
(25G) include, for example, the following compounds:

- 10 2-fluorobromobenzene, 2,6-difluorobromobenzene,
2,4-difluorobromobenzene, 2,3-difluorobromobenzene,
2,4,6-trifluorobromobenzene, 2,4,5-trifluorobromobenzene,
2,3,5,6-pentafluorobromobenzene, tetrafluorobromobenzene,
2-fluoro-5-methylbromobenzene,
15 2-fluoro-4-methylbromobenzene,
2-fluoro-4,6-dimethylbromobenzene,
2-fluoro-4,6-di-tert-butylbromobenzene,
2-fluoro-4-methyl-6-tert-butylbromobenzene,
2-fluoro-6-methyl-4-tert-butylbromobenzene,
20 2-fluoro-5-phenylbromobenzene,
2-fluoro-4-phenylbromobenzene,
2-fluoro-4,6-diphenylbromobenzene,
2-fluoro-5-benzylbromobenzene,
2-fluoro-4-benzylbromobenzene,
25 2-fluoro-4,6-dibenzylbromobenzene,
2-fluoro-5-methoxybromobenzene,
2-fluoro-4-methoxybromobenzene,
2-fluoro-4,6-dimethoxybromobenzene,
2-fluoro-5-aminobromobenzene, 2-fluoro-4-aminobromobenzene,

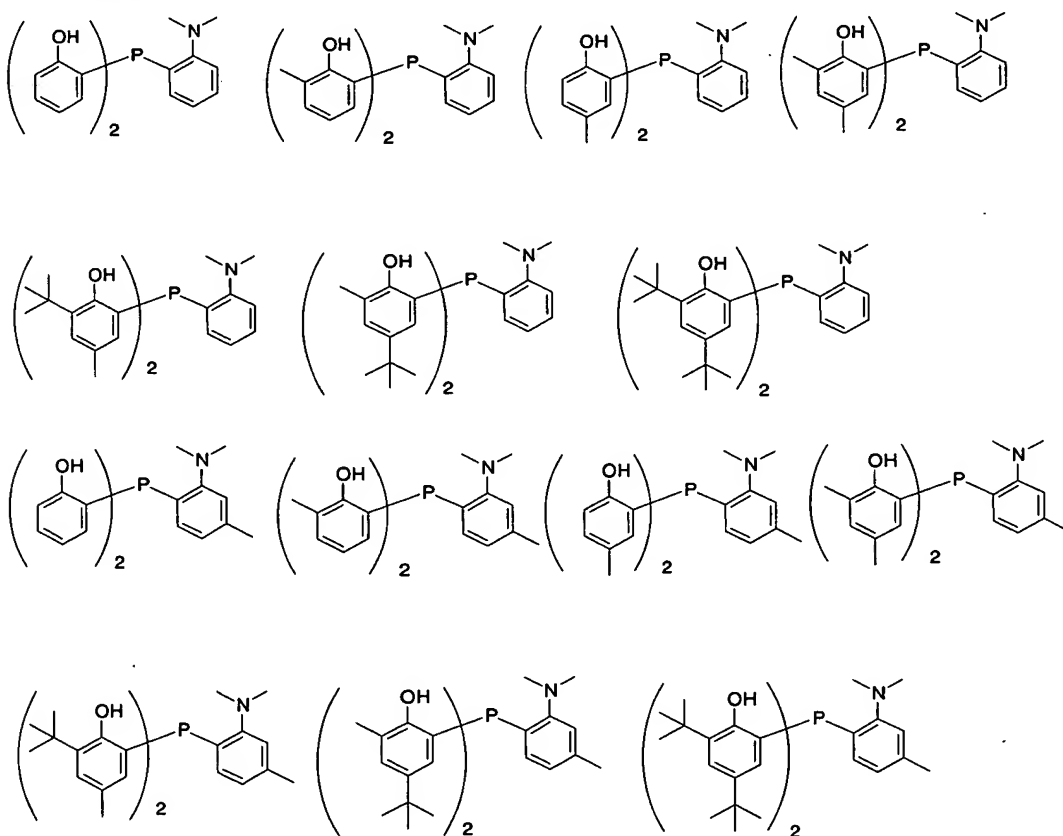
2-fluoro-4,6-diaminobromobenzene,
 2-fluoro-5-(dimethylamono)bromobenzene,
 2-fluoro-4-(dimethylamono)bromobenzene,
 2-fluoro-4,6-bis(dimethylamino)bromobenzene,
 5 2-fluoro-5-(trimethylsilyl)bromobenzene,
 2-fluoro-4-(trimethylsilyl)bromobenzene,
 2-fluoro-4,6-bis(trimethylsilyl)bromobenzene,
 2-trifluoromethylbromobenzene,
 2,6-bis(trifluoromethyl)bromobenzene,
 10 2,4-bis(trifluoromethyl)bromobenzene,
 2,3-bis(trifluoromethyl)bromobenzene,
 2,4,6-tris(trifluoromethyl)bromobenzene,
 2,4,5-tris(trifluoromethyl)bromobenzene,
 2-trifluoromethyl-5-methylbromobenzene,
 15 2-trifluoromethyl-4-methylbromobenzene,
 2-trifluoromethyl-4,6-dimethylbromobenzene,
 2-trifluoromethyl-4,6-di-tert-butylbromobenzene,
 2-trifluoromethyl-4-methyl-6-tert-butylbromobenzene,
 2-trifluoromethyl-6-methyl-4-tert-butylbromobenzene,
 20 2-trifluoromethyl-5-phenylbromobenzene,
 2-trifluoromethyl-4-phenylbromobenzene,
 2-trifluoromethyl-4,6-diphenylbromobenzene,
 2-trifluoromethyl-5-benzylbromobenzene,
 2-trifluoromethyl-4-benzylbromobenzene,
 25 2-trifluoromethyl-4,6-dibenzylbromobenzene,
 2-trifluoromethyl-5-methoxybromobenzene,
 2-trifluoromethyl-4-methoxybromobenzene,
 2-trifluoromethyl-4,6-dimethoxybromobenzene,
 2-trifluoromethyl-5-aminobromobenzene,

2-trifluoromethyl-4-aminobromobenzene,
 2-trifluoromethyl-4,6-diaminobromobenzene,
 2-trifluoromethyl-5-(dimethylamino)bromobenzene,
 2-trifluoromethyl-4-(dimethylamino)bromobenzene,
 5 2-trifluoromethyl-4,6-bis(dimethylamino)bromobenzene,
 2-trifluoromethyl-5-(trimethylsilyl)bromobenzene,
 2-trifluoromethyl-4-(trimethylsilyl)bromobenzene,
 2-trifluoromethyl-4,6-bis(trimethylsilyl)bromobenzene,
 2-fluorochlorobenzene, 2,6-difluorochlorobenzene,
 10 2,4-difluorochlorobenzene, 2,3-difluorochlorobenzene,
 2,4,6-trifluorochlorobenzene, 2,4,5-trifluorochlorobenzene,
 2,3,5,6-pentafluorochlorobenzene, tetrafluorochlorobenzene,
 2-fluoro-5-methylchlorobenzene,
 2-fluoro-4-methylchlorobenzene,
 15 2-fluoro-4,6-dimethylchlorobenzene,
 2-fluoro-4,6-di-tert-butylchlorobenzene,
 2-fluoro-4-methyl-6-tert-butylchlorobenzene,
 2-fluoro-6-methyl-4-tert-butylchlorobenzene,
 2-fluoro-5-phenylchlorobenzene,
 20 2-fluoro-4-phenylchlorobenzene,
 2-fluoro-4,6-diphenylchlorobenzene,
 2-fluoro-5-benzylchlorobenzene,
 2-fluoro-4-benzylchlorobenzene,
 2-fluoro-4,6-dibenzylchlorobenzene,
 25 2-fluoro-5-methoxychlorobenzene,
 2-fluoro-4-methoxychlorobenzene,
 2-fluoro-4,6-dimethoxychlorobenzene,
 2-fluoro-5-aminochlorobenzene,
 2-fluoro-4-aminochlorobenzene,

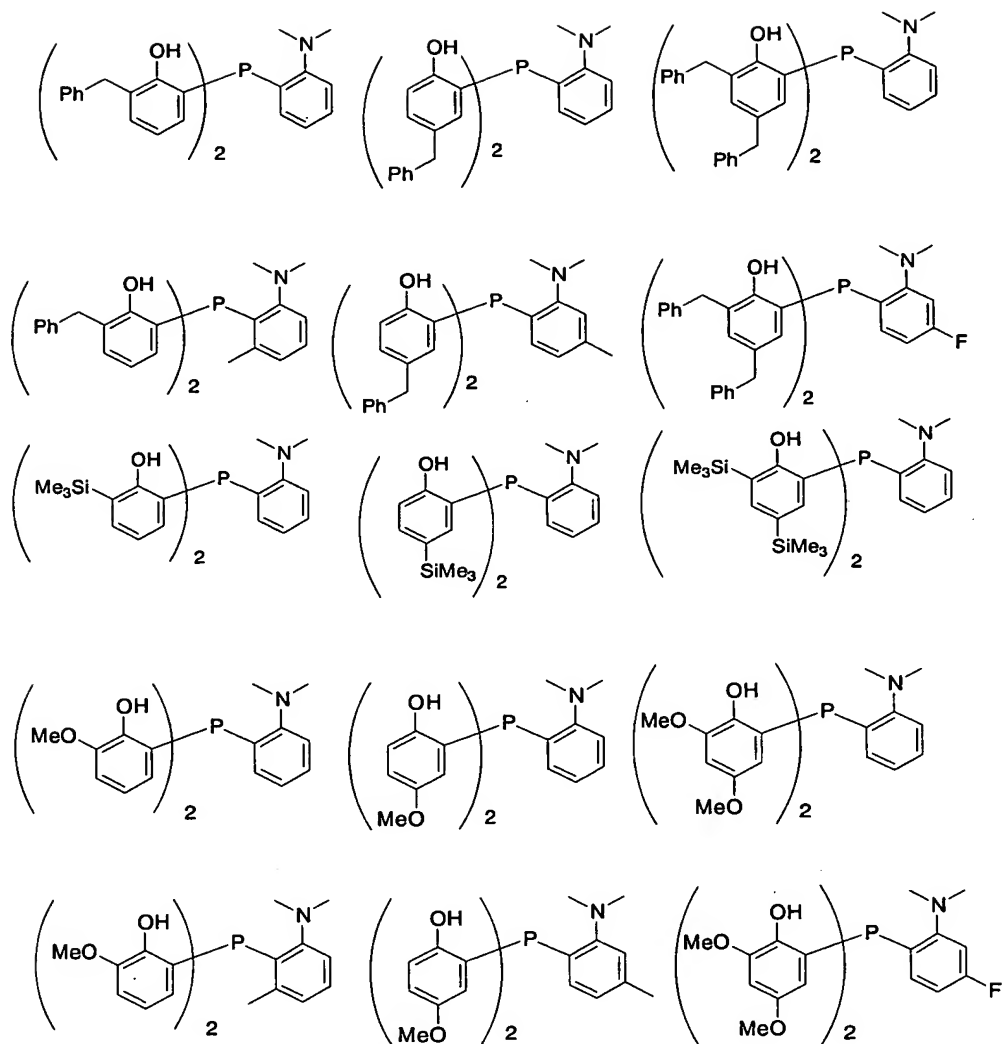
2-fluoro-4,6-diaminochlorobenzene,
 2-fluoro-5-(dimethylamino)chlorobenzene,
 2-fluoro-4-(dimethylamino)chlorobenzene,
 2-fluoro-4,6-bis(dimethylamino)chlorobenzene,
 5 2-fluoro-5-(trimethylsilyl)chlorobenzene,
 2-fluoro-4-(trimethylsilyl)chlorobenzene,
 2-fluoro-4,6-bis(trimethylsilyl)chlorobenzene,
 2-trifluoromethylchlorobenzene,
 2,6-bis(trifluoromethyl)chlorobenzene,
 10 2,4-bis(trifluoromethyl)chlorobenzene,
 2,3-bis(trifluoromethyl)chlorobenzene,
 2,4,6-tris(trifluoromethyl)chlorobenzene,
 2,4,5-tris(trifluoromethyl)chlorobenzene,
 2-trifluoromethyl-5-methylchlorobenzene,
 15 2-trifluoromethyl-4-methylchlorobenzene,
 2-trifluoromethyl-4,6-dimethylchlorobenzene,
 2-trifluoromethyl-4,6-di-tert-butylchlorobenzene,
 2-trifluoromethyl-4-methyl-6-tert-butylchlorobenzene,
 2-trifluoromethyl-6-methyl-4-tert-butylchlorobenzene,
 20 2-trifluoromethyl-5-phenylchlorobenzene,
 2-trifluoromethyl-4-phenylchlorobenzene,
 2-trifluoromethyl-4,6-diphenylchlorobenzene,
 2-trifluoromethyl-5-benzylchlorobenzene,
 2-trifluoromethyl-4-benzylchlorobenzene,
 25 2-trifluoromethyl-4,6-dibenzylchlorobenzene,
 2-trifluoromethyl-5-methoxychlorobenzene,
 2-trifluoromethyl-4-methoxychlorobenzene,
 2-trifluoromethyl-4,6-dimethoxychlorobenzene,
 2-trifluoromethyl-5-aminochlorobenzene,

- 2-trifluoromethyl-4-aminochlorobenzene,
 2-trifluoromethyl-4,6-diaminochlorobenzene,
 2-trifluoromethyl-5-(dimethylamino)chlorobenzene,
 2-trifluoromethyl-4-(diaminomethyl)chlorobenzene,
 5 2-trifluoromethyl-4,6-bis(dimethylamino)chlorobenzene,
 2-trifluoromethyl-5-(trimethylsilyl)chlorobenzene,
 2-trifluoromethyl-4-(trimethylsilyl)chlorobenzene, and
 2-trifluoromethyl-4,6-bis(trimethylsilyl)chlorobenzene.

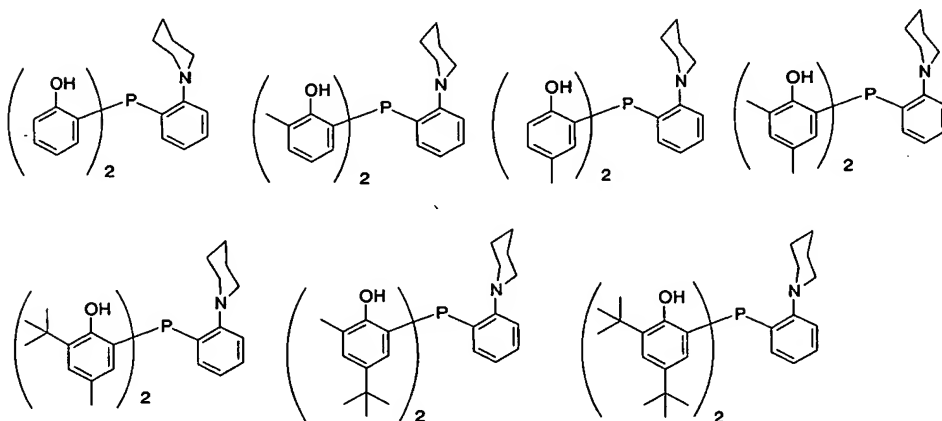
Examples of the compound of formula (1) wherein G^2 is G^{26} ,
 10 which corresponds to the compound of formula (26A) include, the
 following compounds:

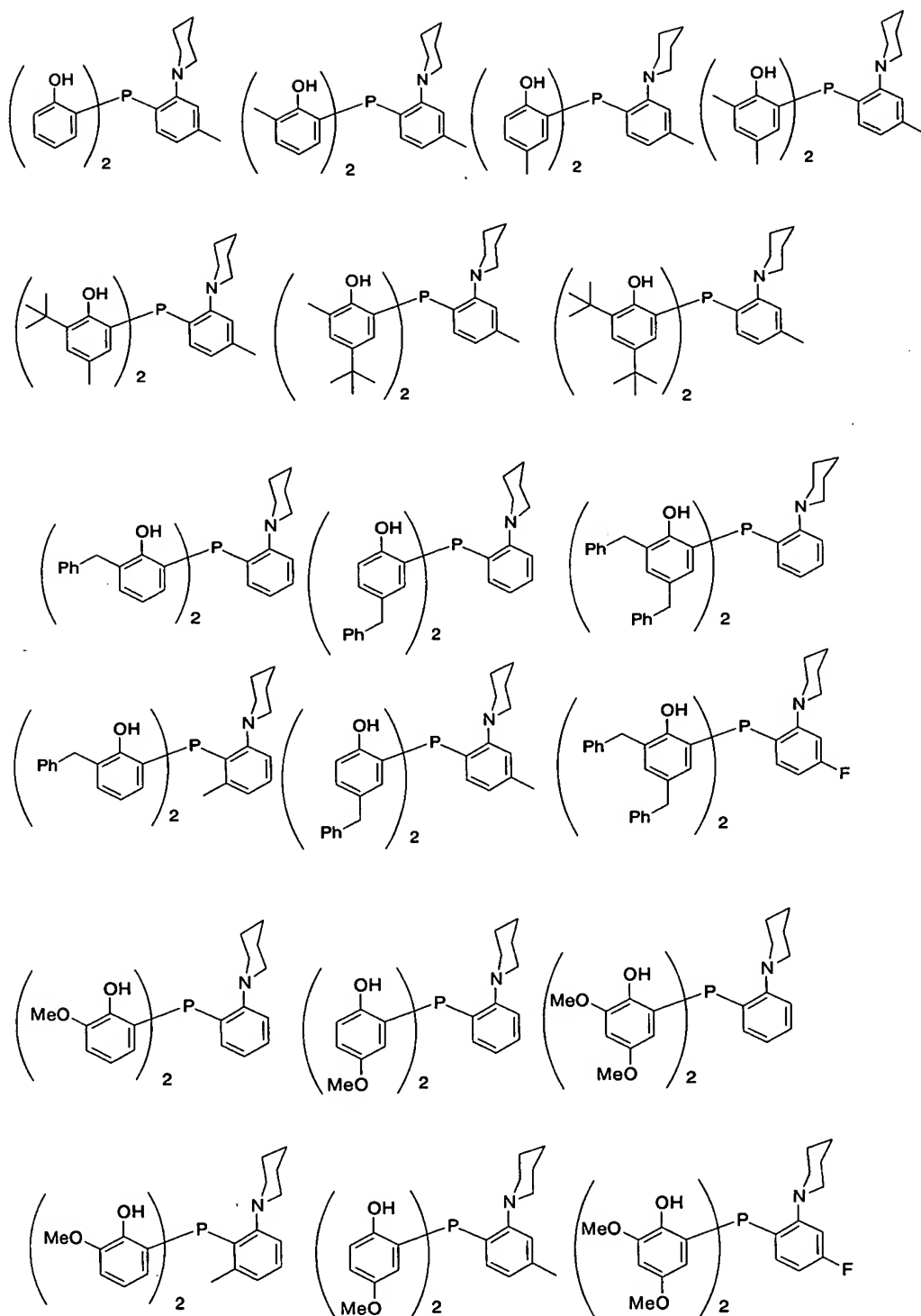


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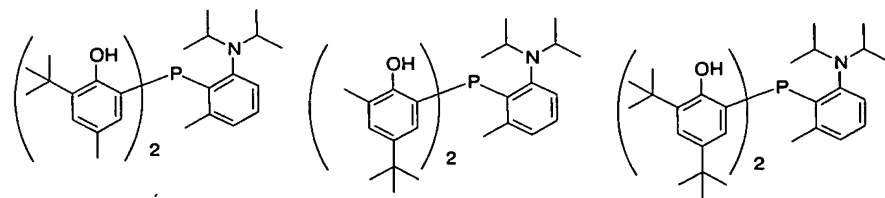
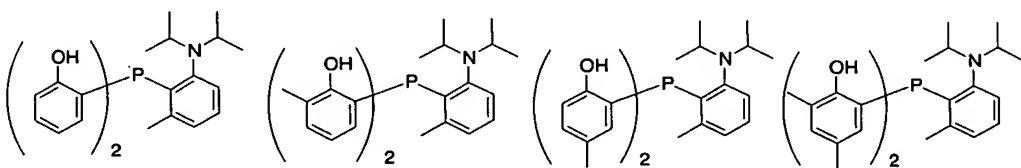
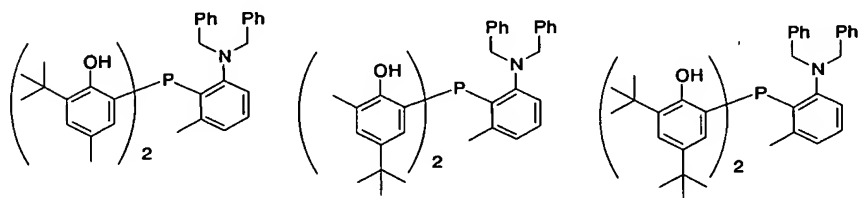
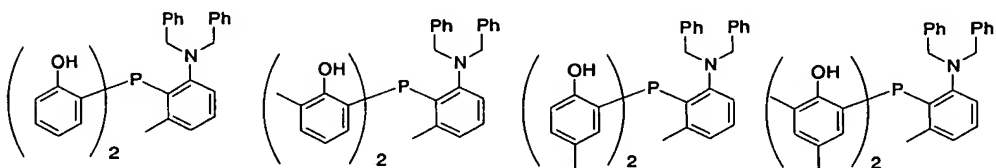
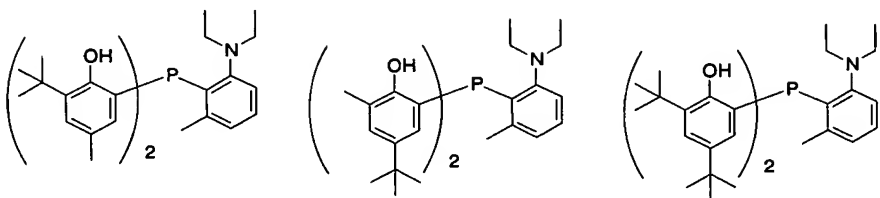
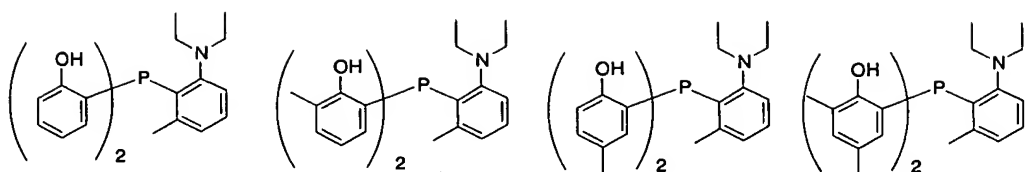


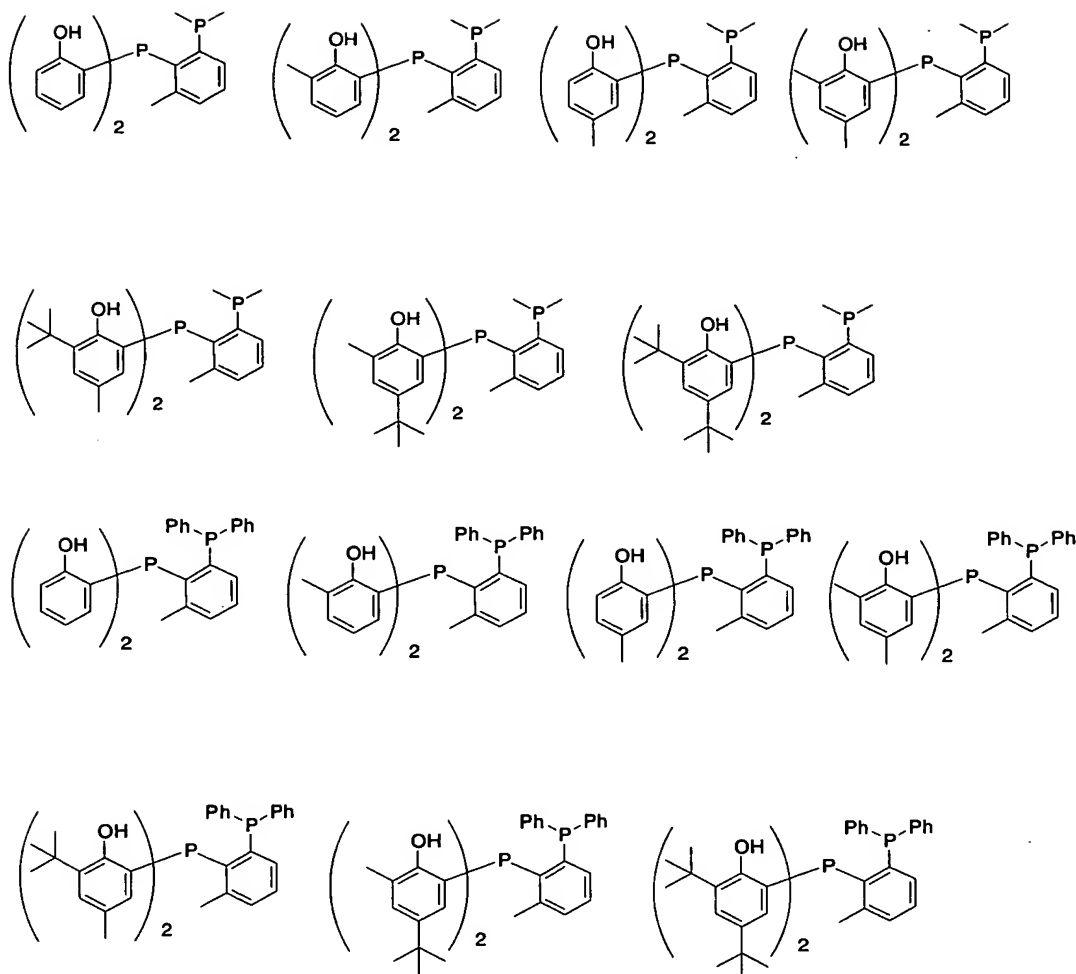
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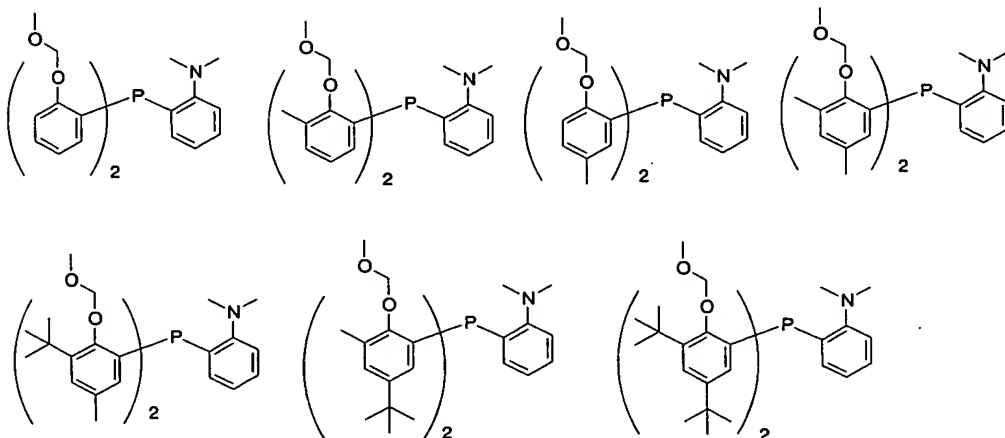


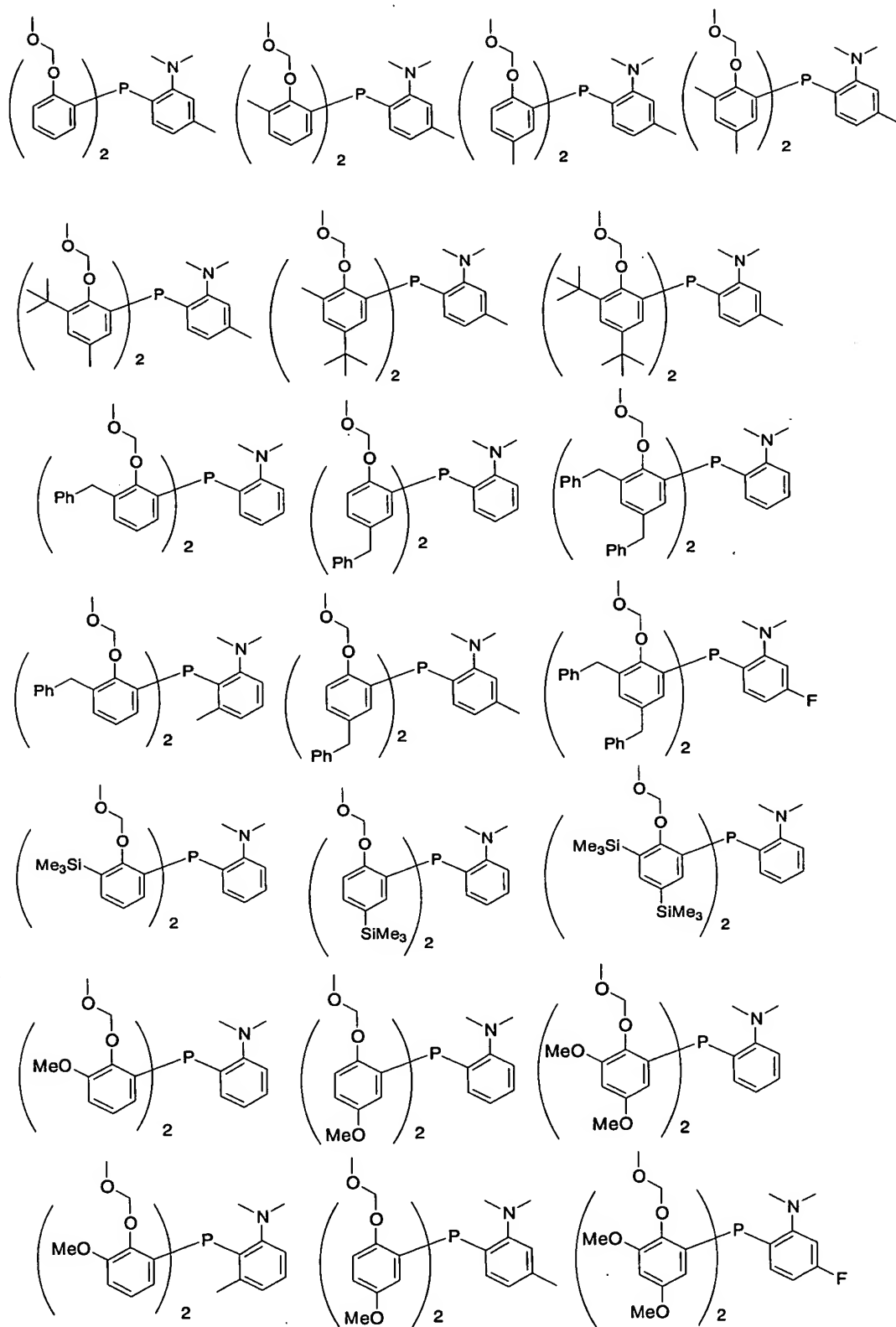
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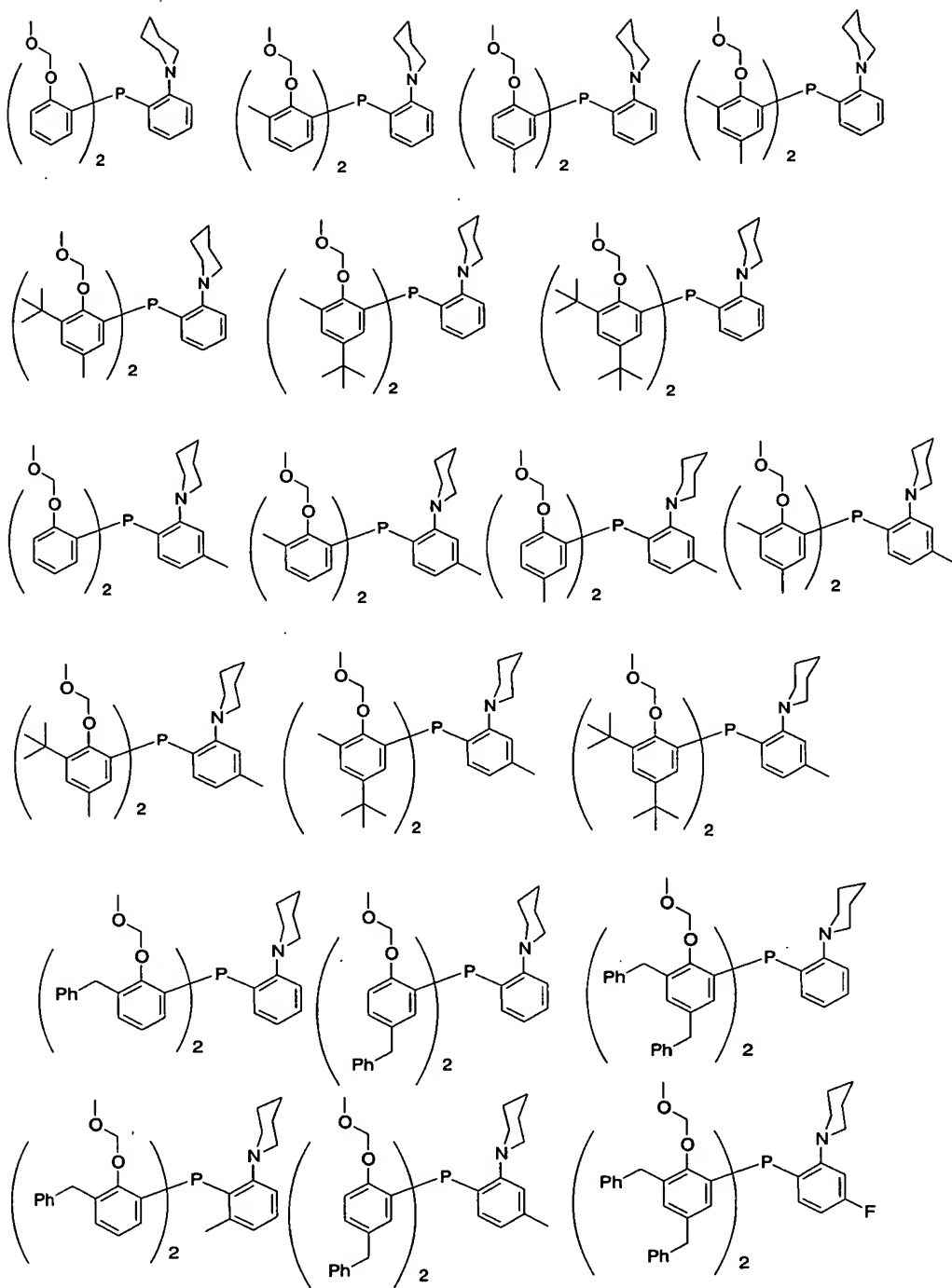




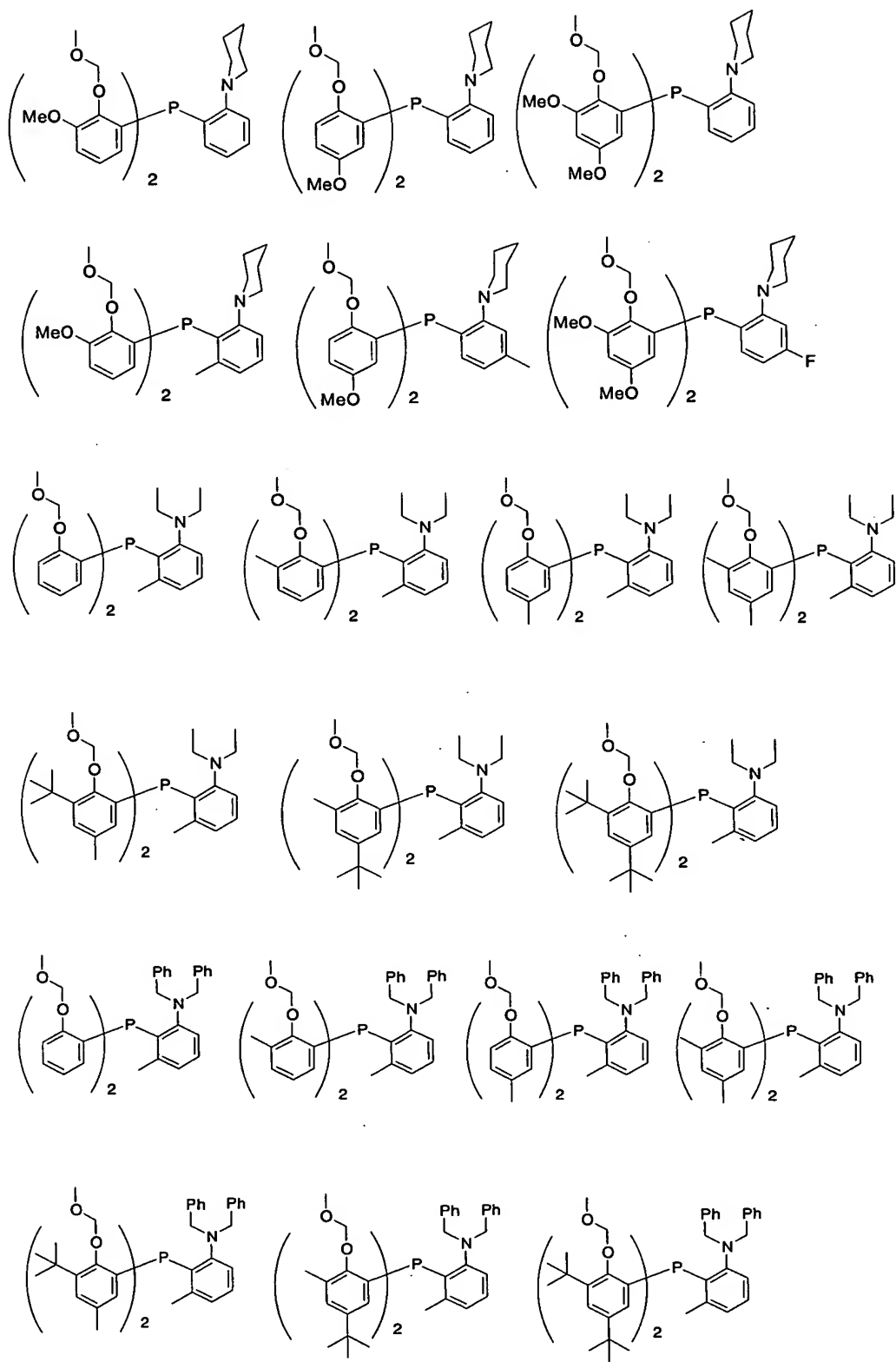
5 Examples of the compound of formula (1) wherein G^2 in formula 1 is G^{26} , which corresponds to the compound of formula (26B) include, for example, the following compounds:

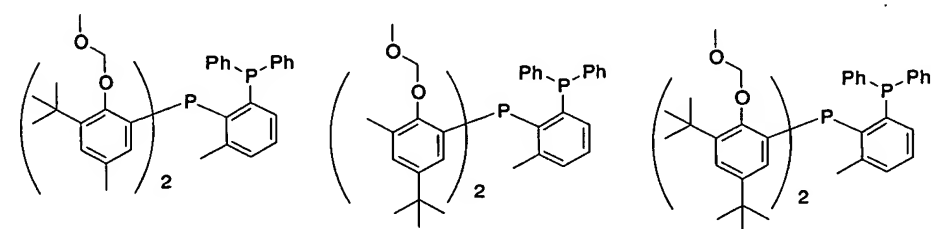
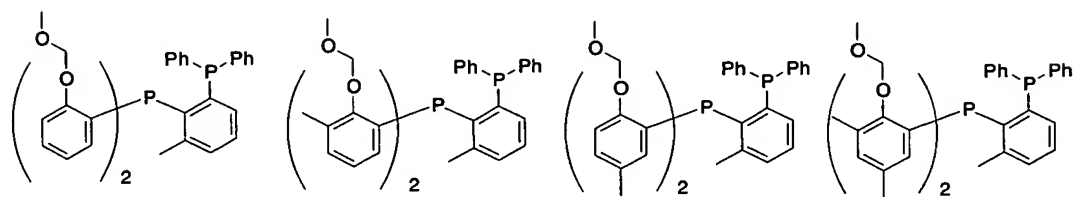
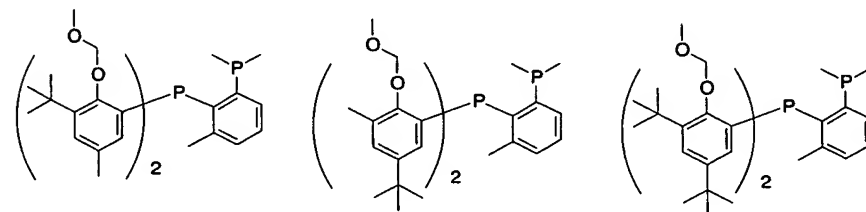
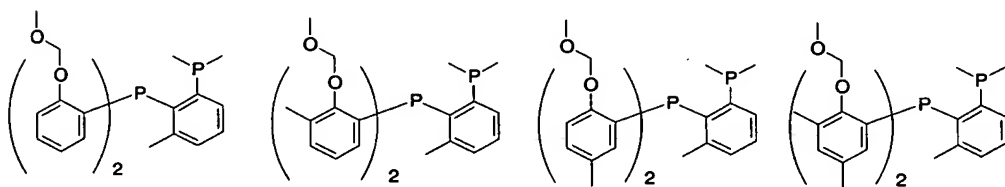
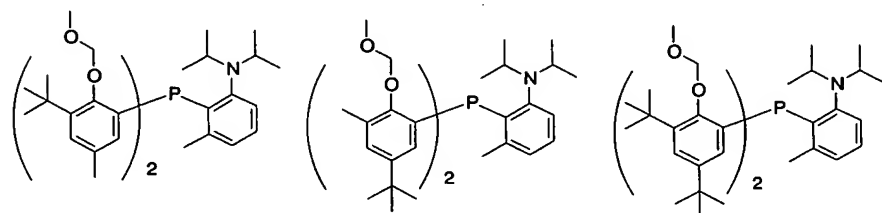
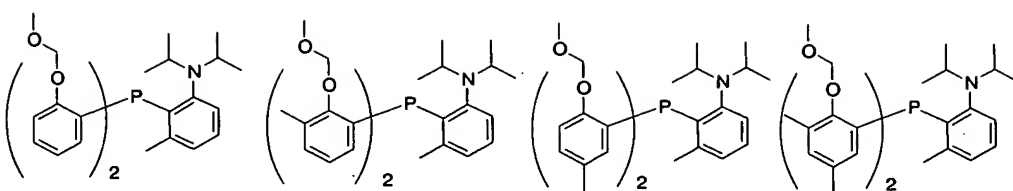


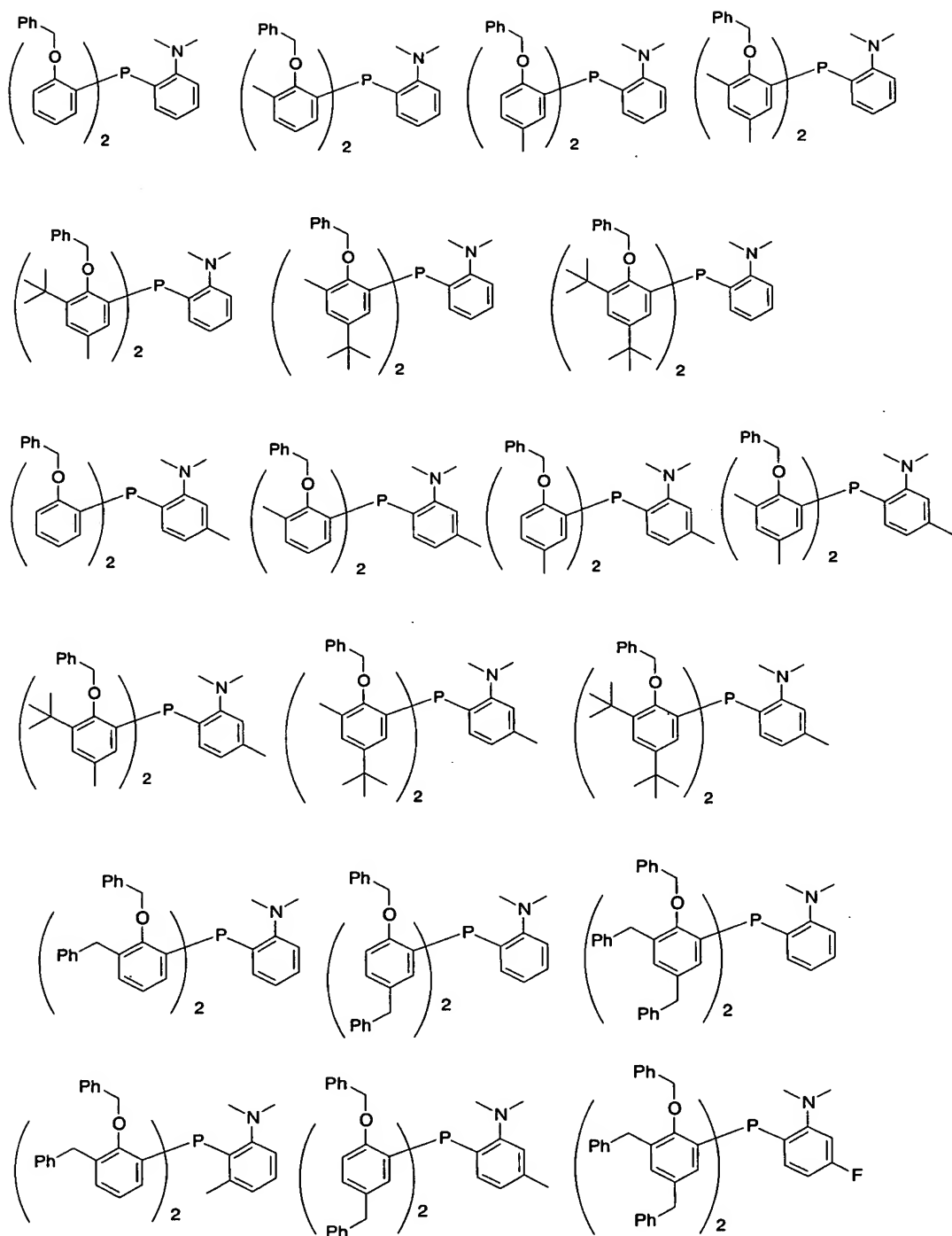




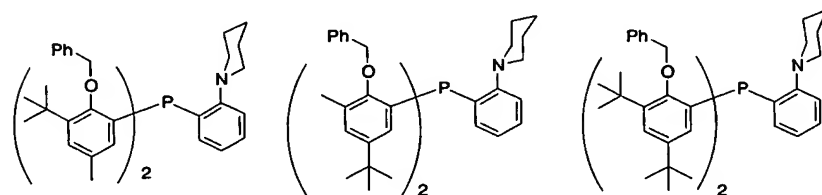
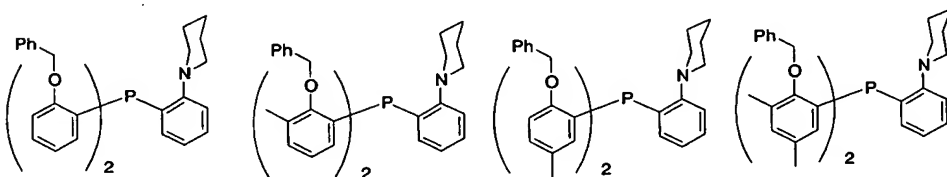
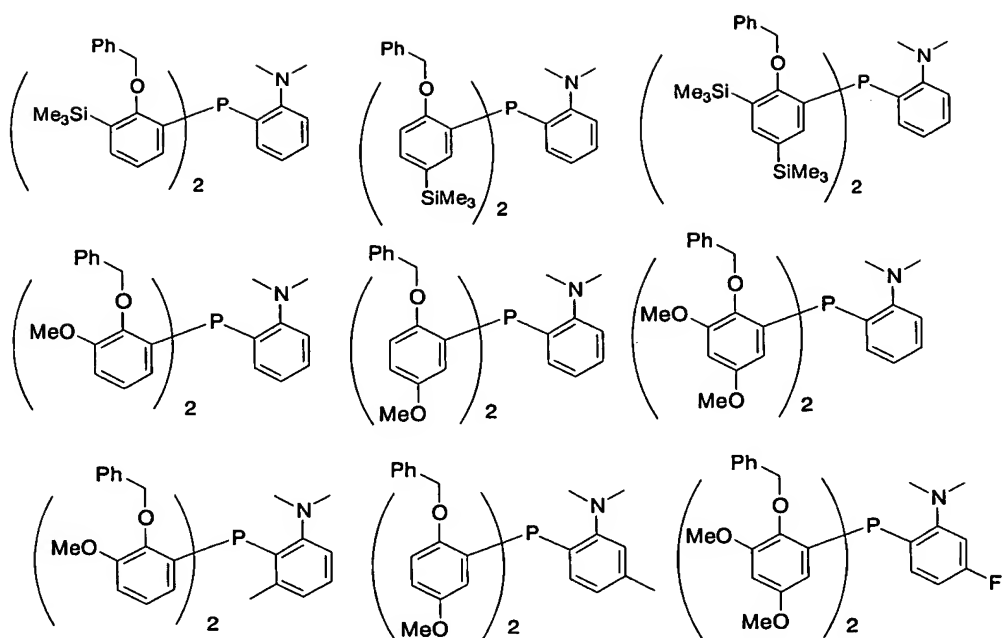
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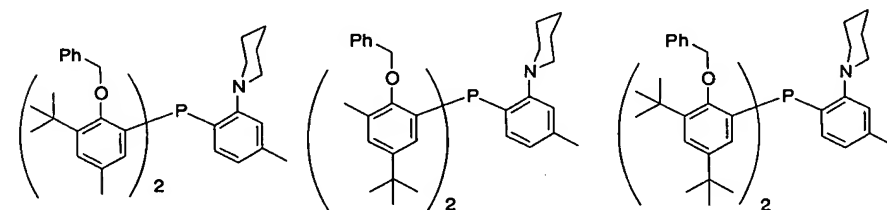
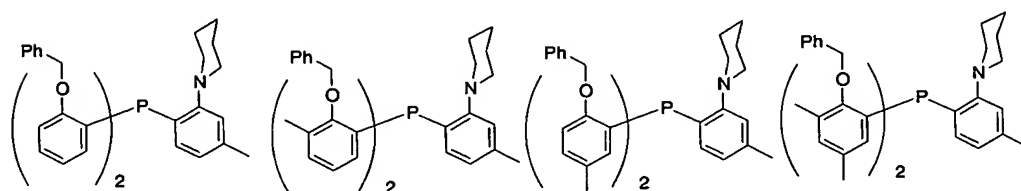


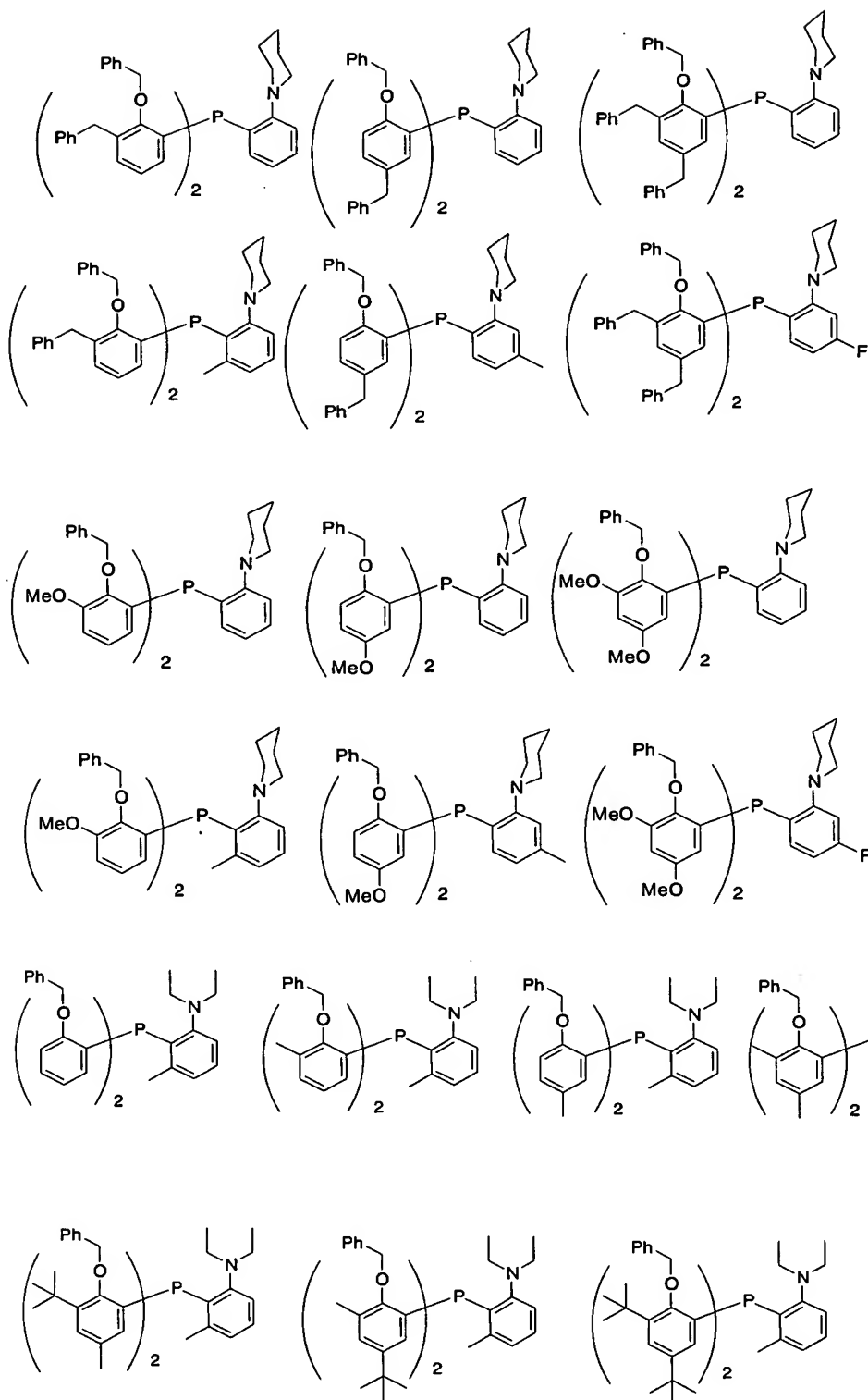


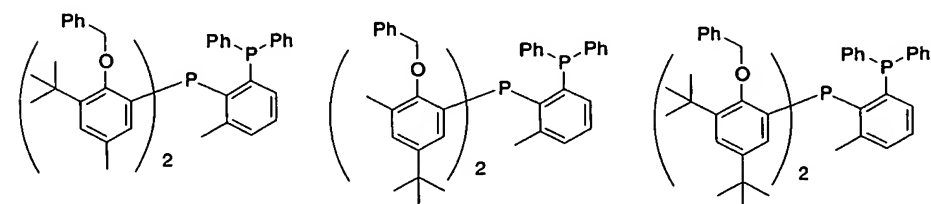
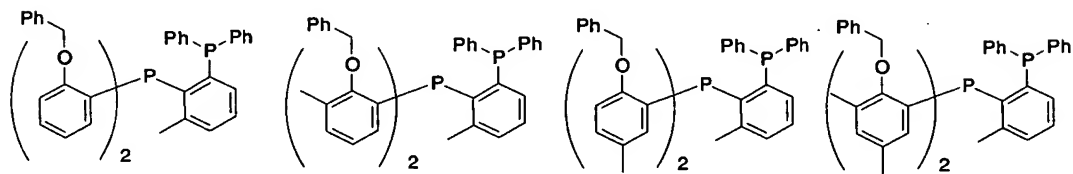
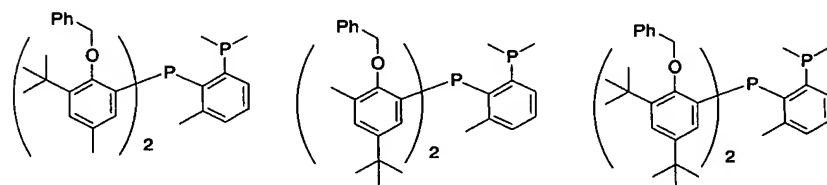
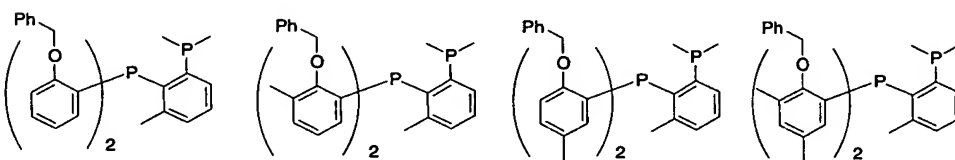
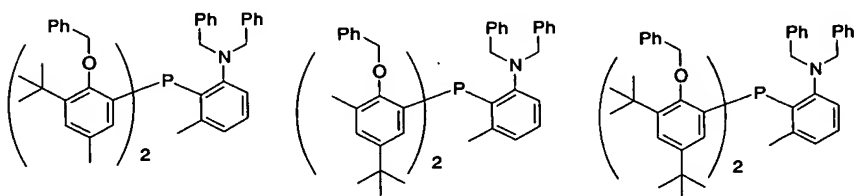
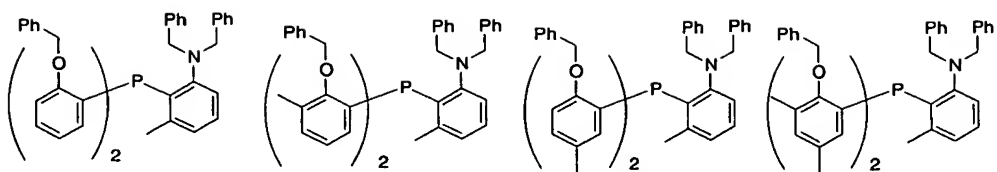
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5

The phosphine compound of formula (26B) can be produced by reacting phosphine dihalide of formula (26C) with the metal

aryl compound of formula (26D).

The molar ratio between phosphine dihalide of formula (26C) and the metal aryl compound of formula (26D) is not particularly restricted, and it is preferably in the range of 1:0.5 to 1:5,
5 more preferably 1:1 to 1:2.5.

Preferable examples of the halogen atom represented by X^2 in formula (26C) or (26D) include chlorine, bromine and iodine atoms, and the chlorine atom is preferable.

Specific examples of the alkali metal and alkaline earth
10 metal represented by D in formula (26D) include lithium, sodium, potassium, magnesium and calcium atoms, and the lithium and magnesium atoms are preferable.

The reaction above is usually performed in a solvent inert to the reaction. Examples of the solvent include aromatic
15 hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture of at least two of them. The amount thereof is usually 1 to
20 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (25D).

The reaction can be performed by adding, for example, phosphine dihalide of formula (26C) to the metal aryl compound of formula (26D). The reaction temperature is usually in the
25 range of from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C .

The phosphine compound of formula (26B) is obtained, for example, by removing insolubles by filtration followed by removing the solvent by evaporation. The product can be purified

by silica gel column chromatography, if necessary.

The phosphine compound of formula (26B) can be produced by reacting phosphine halide of formula (26E) with the aryl compound of formula (26F).

5 The molar ratio between phosphine halide of formula (26E) and the aryl compound of formula (26F) in the reaction is not particularly restricted, and the ratio is preferably in the range of 1:0.1 to 1:5, more preferably 1:0.5 to 1:2.

10 Specific examples of the halogen atom represented by X^2 in formula (26E) or (26F) include fluorine, chlorine, bromine and iodine atoms, and the chlorine atom is preferable.

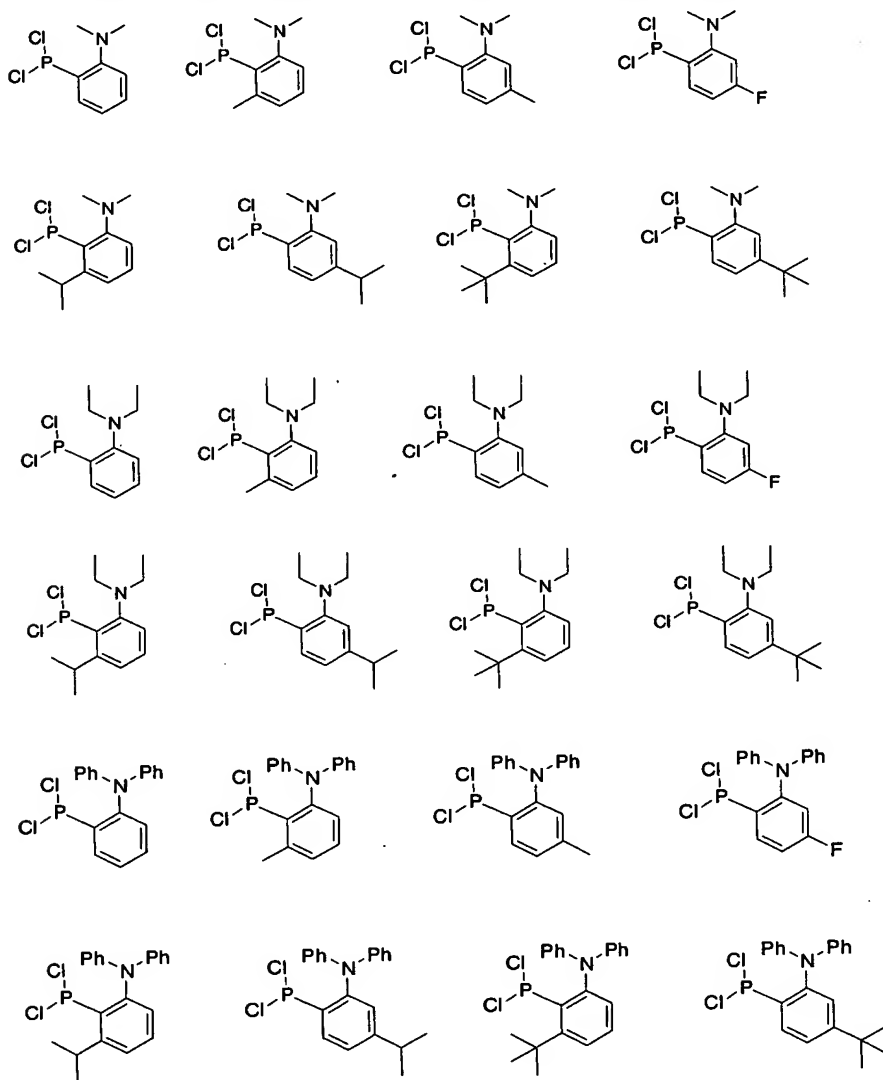
15 Specific examples of the alkali metal and alkaline earth metal represented by D in formula (26F) include lithium, sodium, potassium, magnesium and calcium atoms, and lithium and magnesium atoms are preferable.

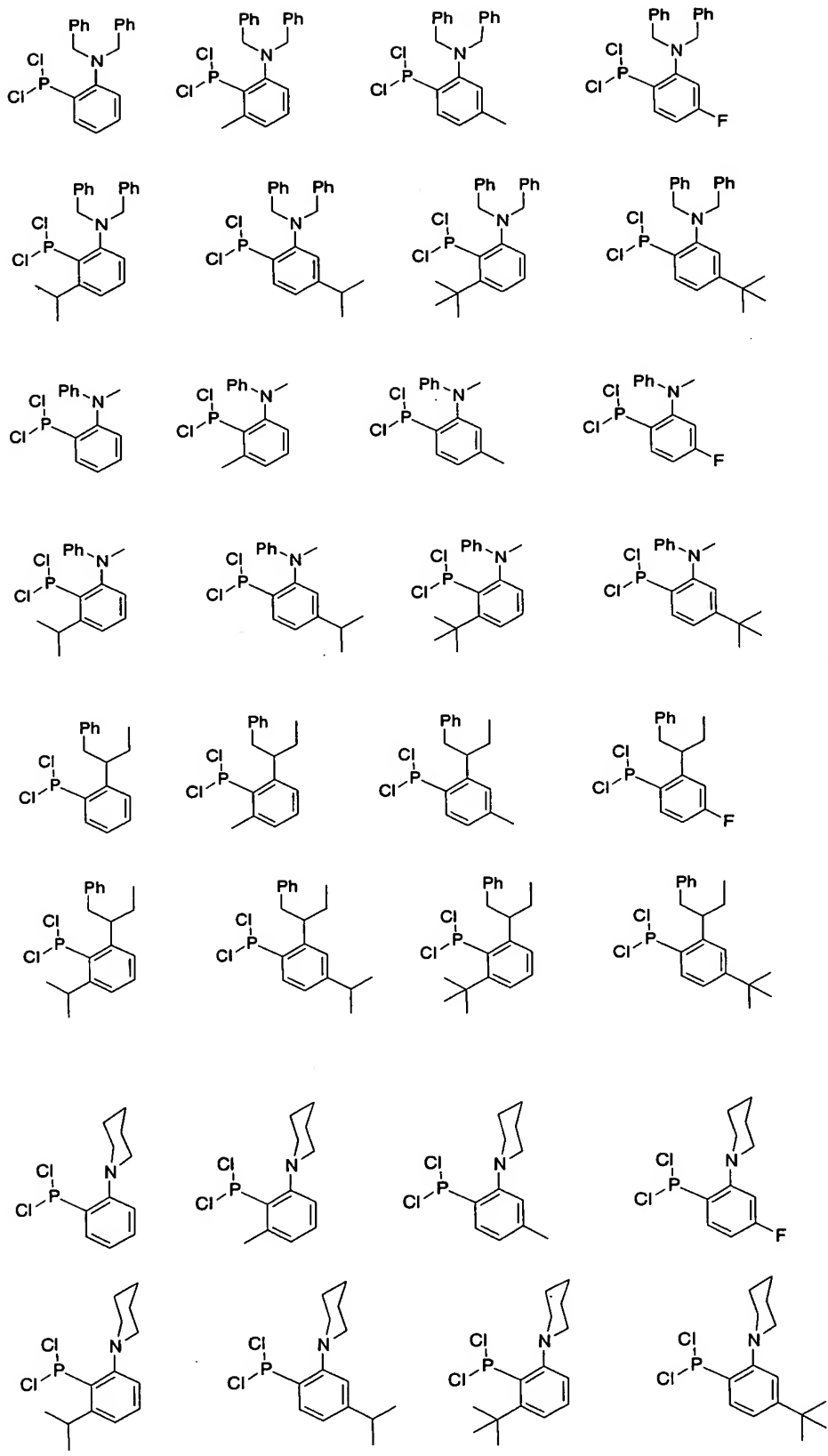
20 The reaction above is usually performed in a solvent inert to the reaction. Examples of the solvent include aromatic hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; and ether solvents such as diethyl ether, tetrahydrofuran or the like. These solvents may be used alone or as a mixture of at least two of them. The amount thereof is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the metal aryl compound of formula (8).

25 The reaction can be performed by adding, for example, phosphine halide of formula 26E to the aryl compound of formula (26F). The reaction temperature is usually in the range of from -100°C or more to the boiling point or less of the solvent, more preferably in the range of -80°C to 100°C.

The phosphine compound of formula (26B) is obtained, for example, by removing insolubles by filtration followed by removing the solvent by evaporation. The product may be purified by silica gel column chromatography, if necessary.

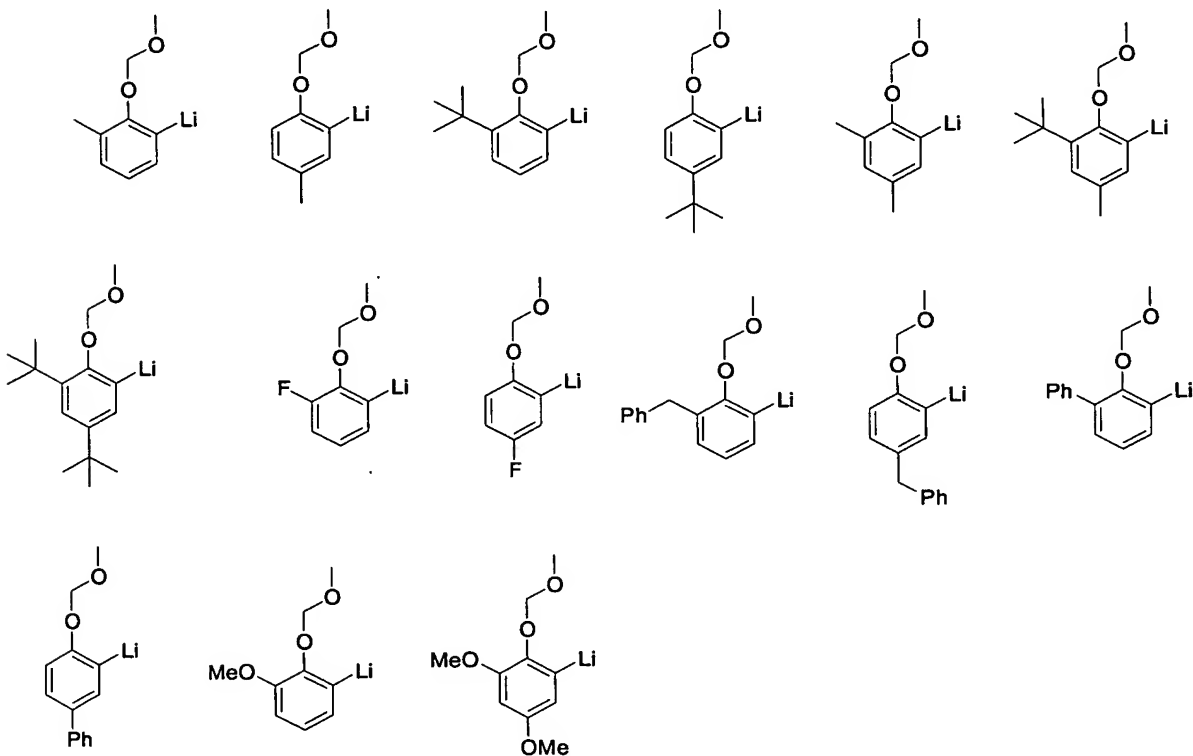
5 Specific examples of phosphine dihalide of formula (26C) include, for example, the following compounds:

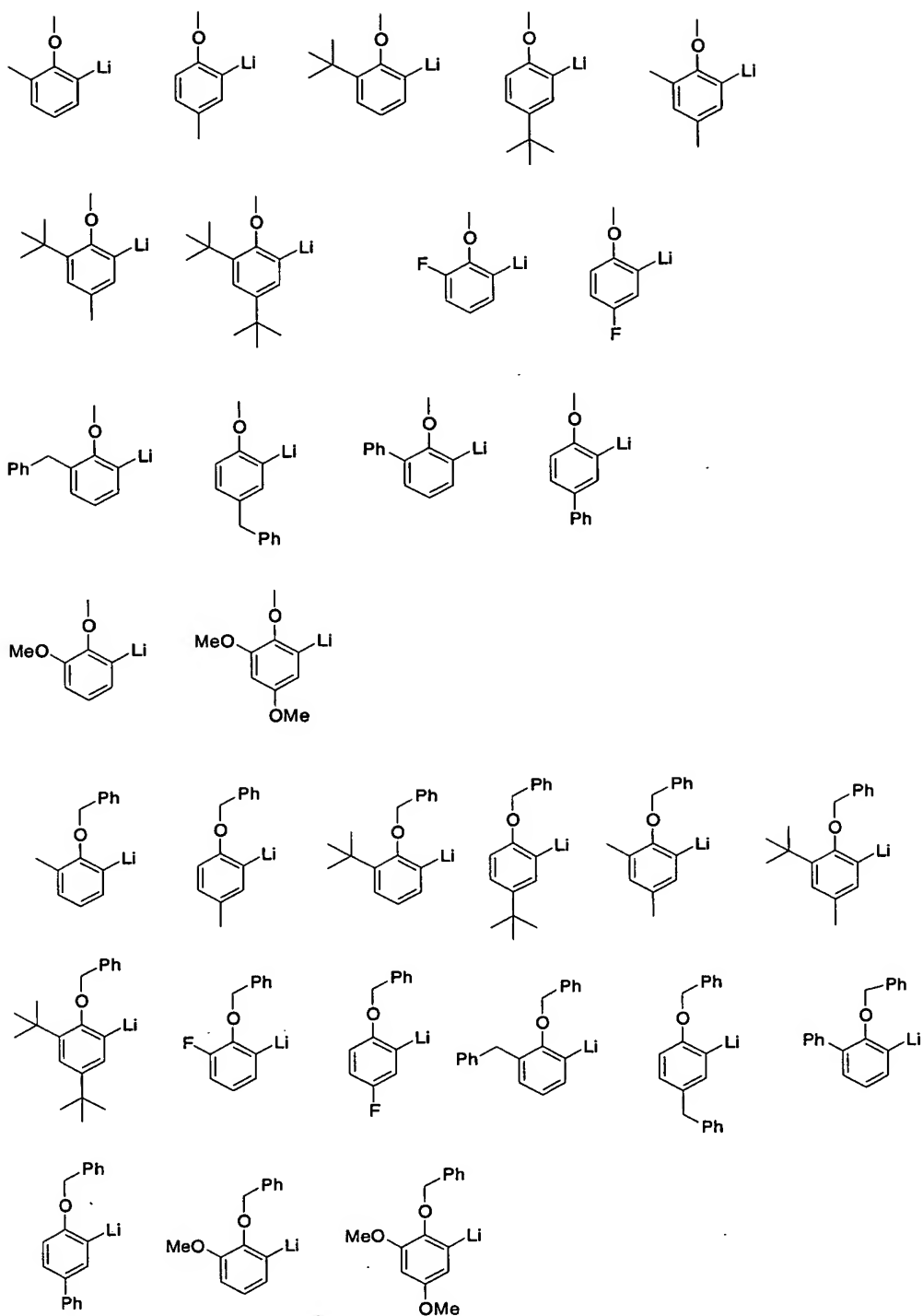


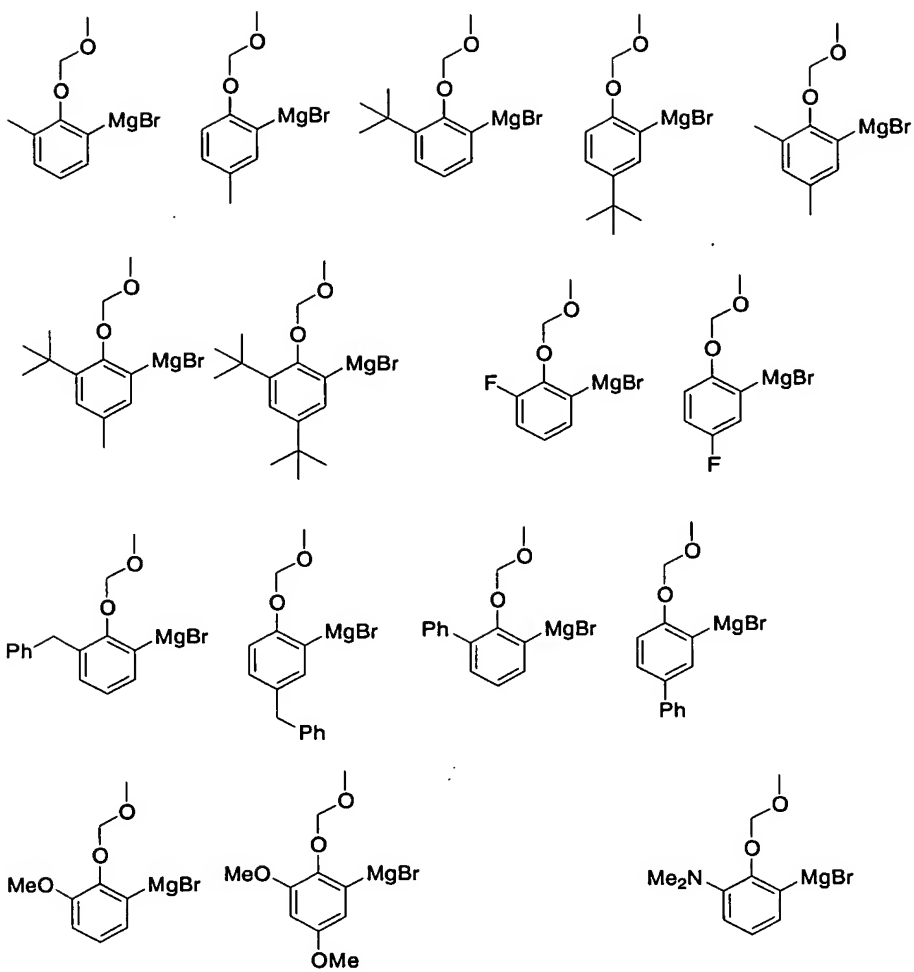


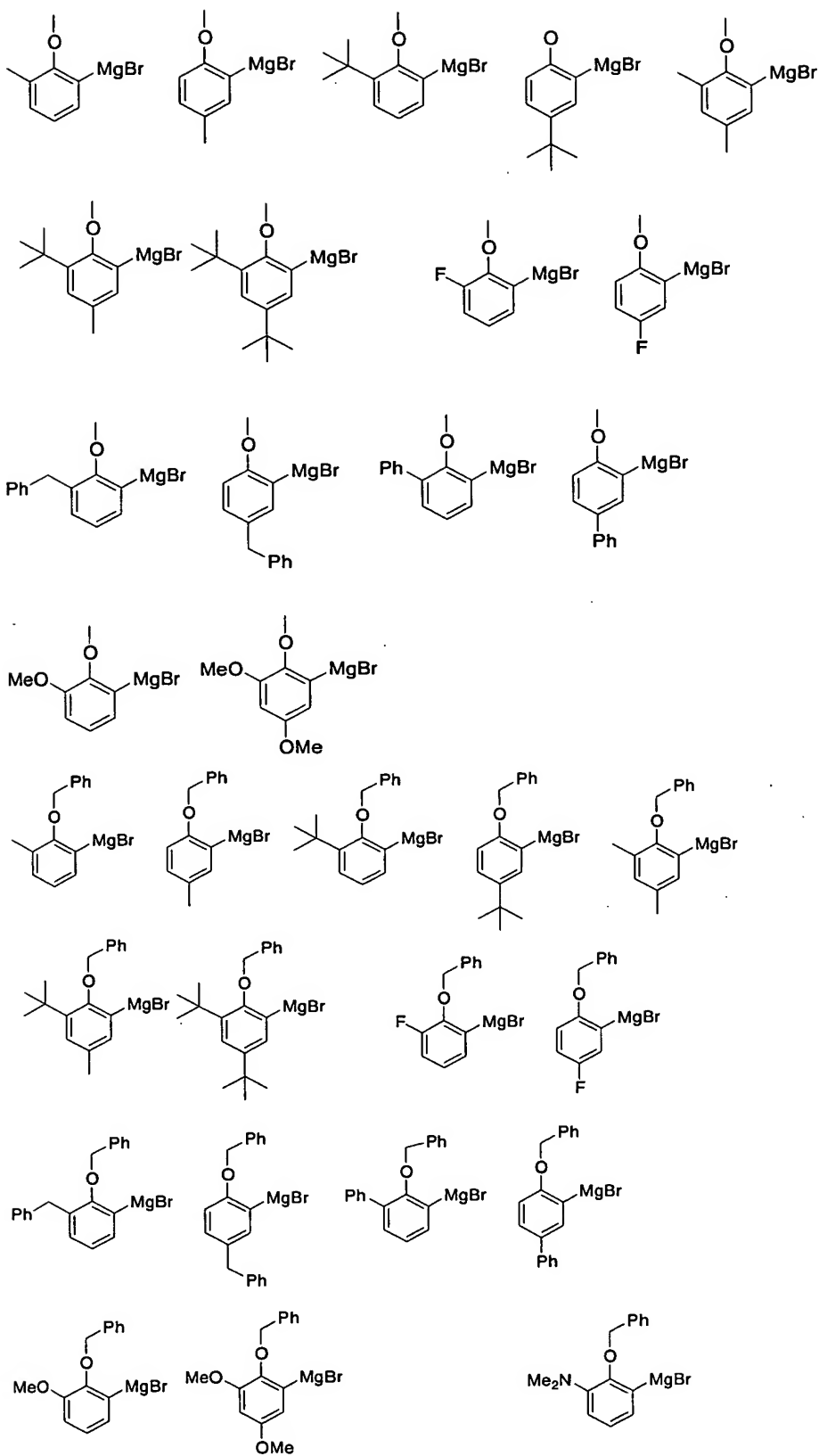
The compounds of formulae above also include those in which the chlorine atom is replaced by bromine or iodine atoms.

Specific examples of the metal aryl compound of formula (26D) include, for example, the following compounds:

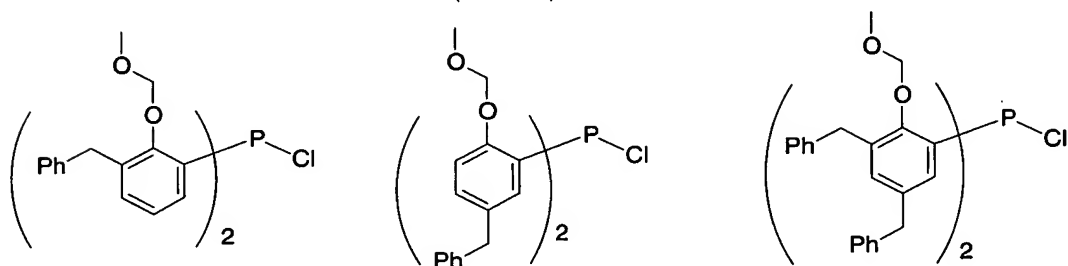
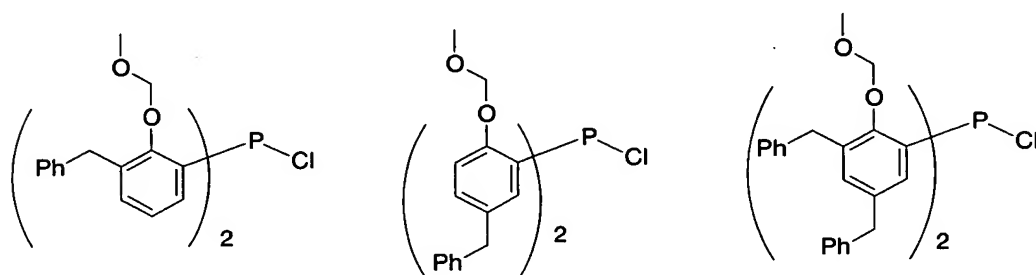
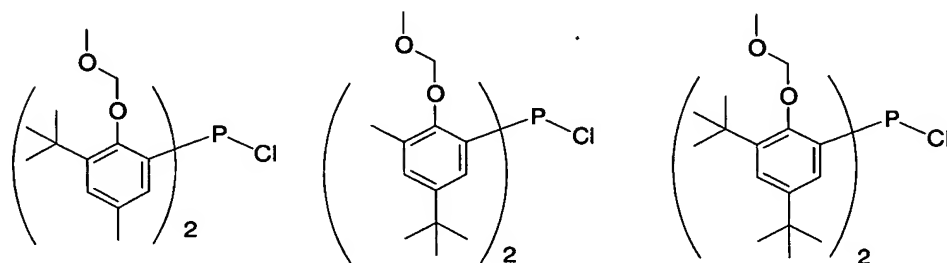
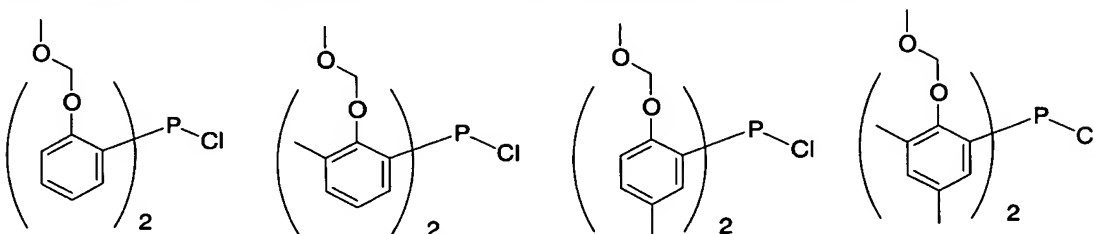




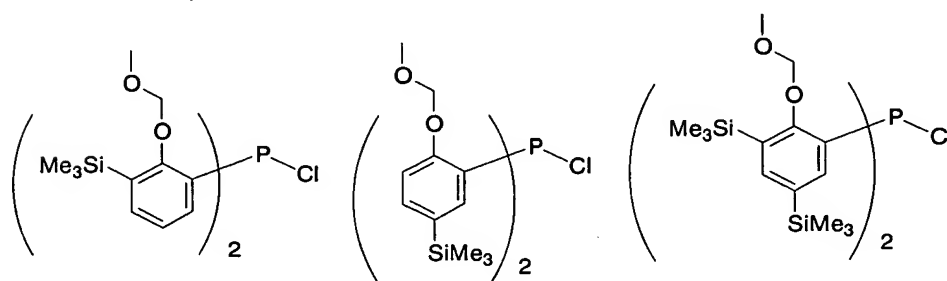


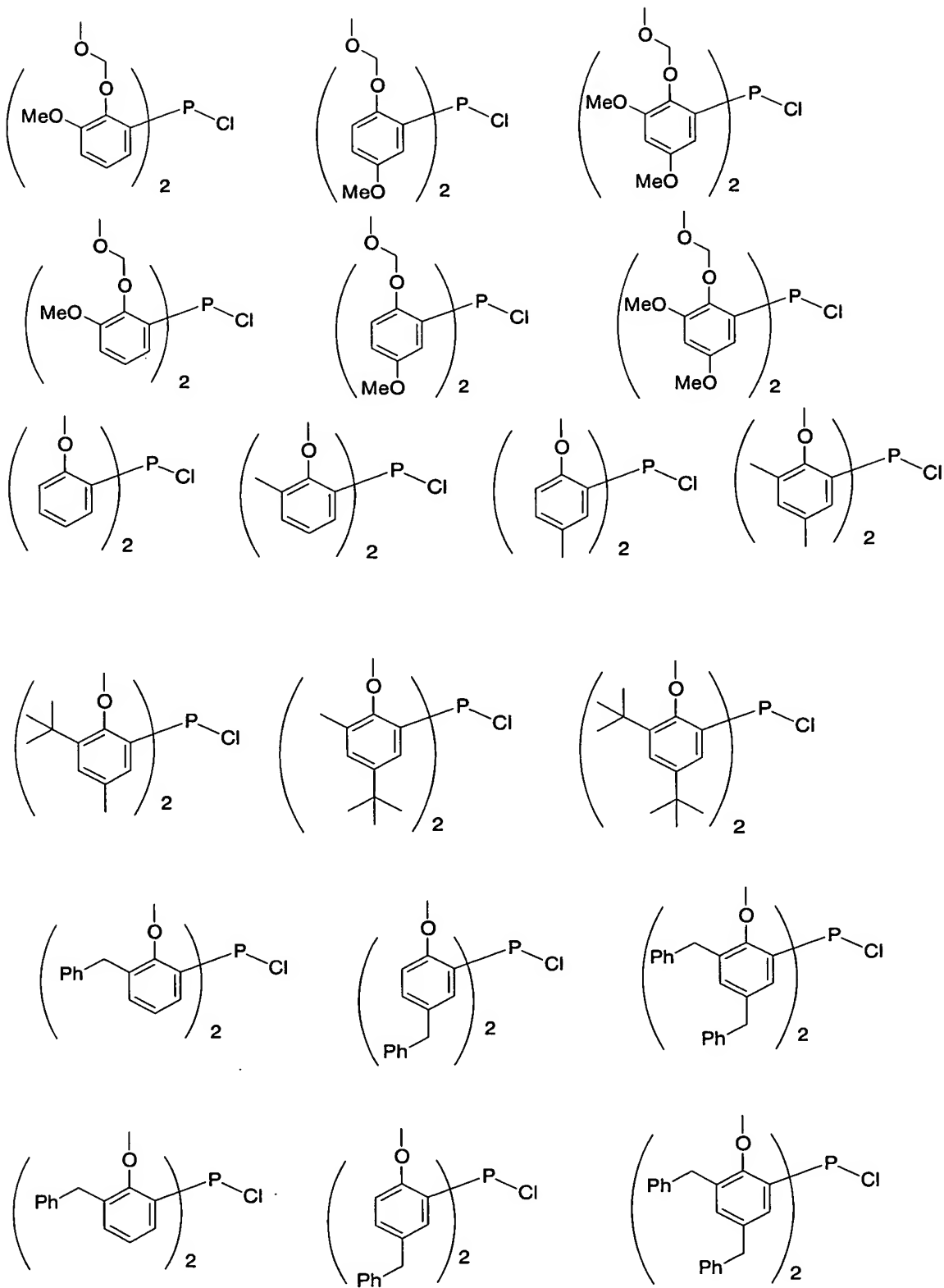


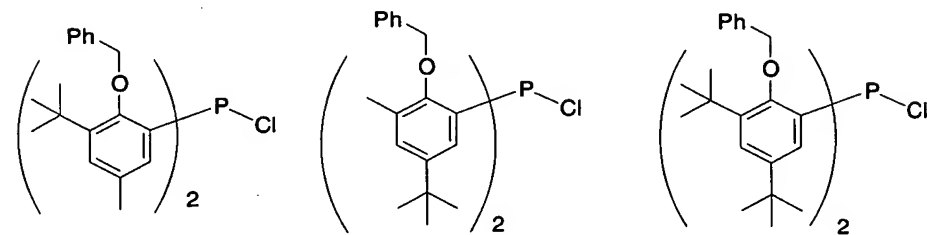
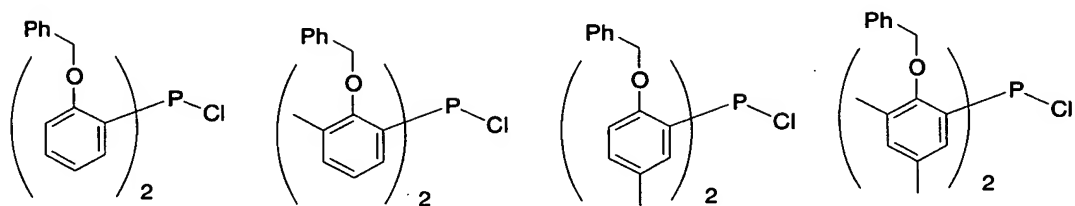
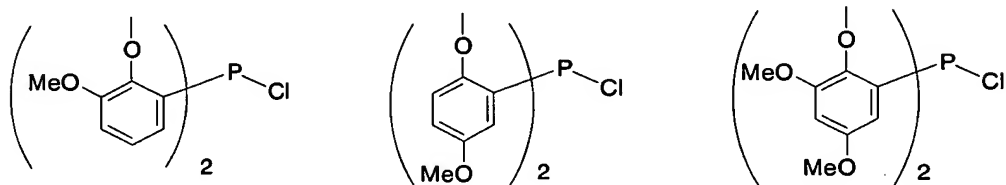
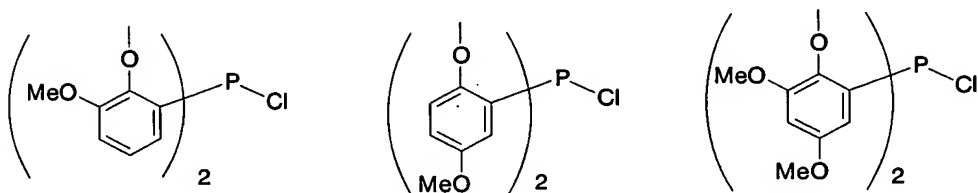
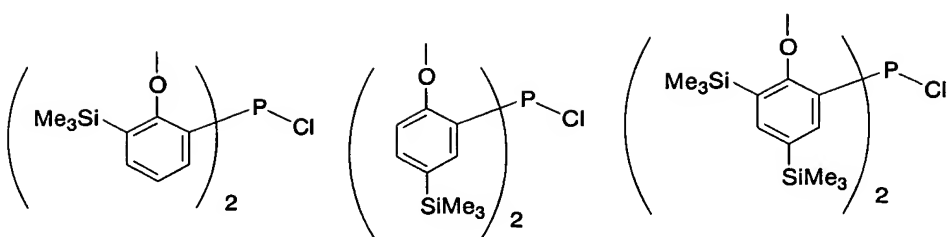
Specific examples of phosphine halide of formula (26E) include, for example, the following compounds:



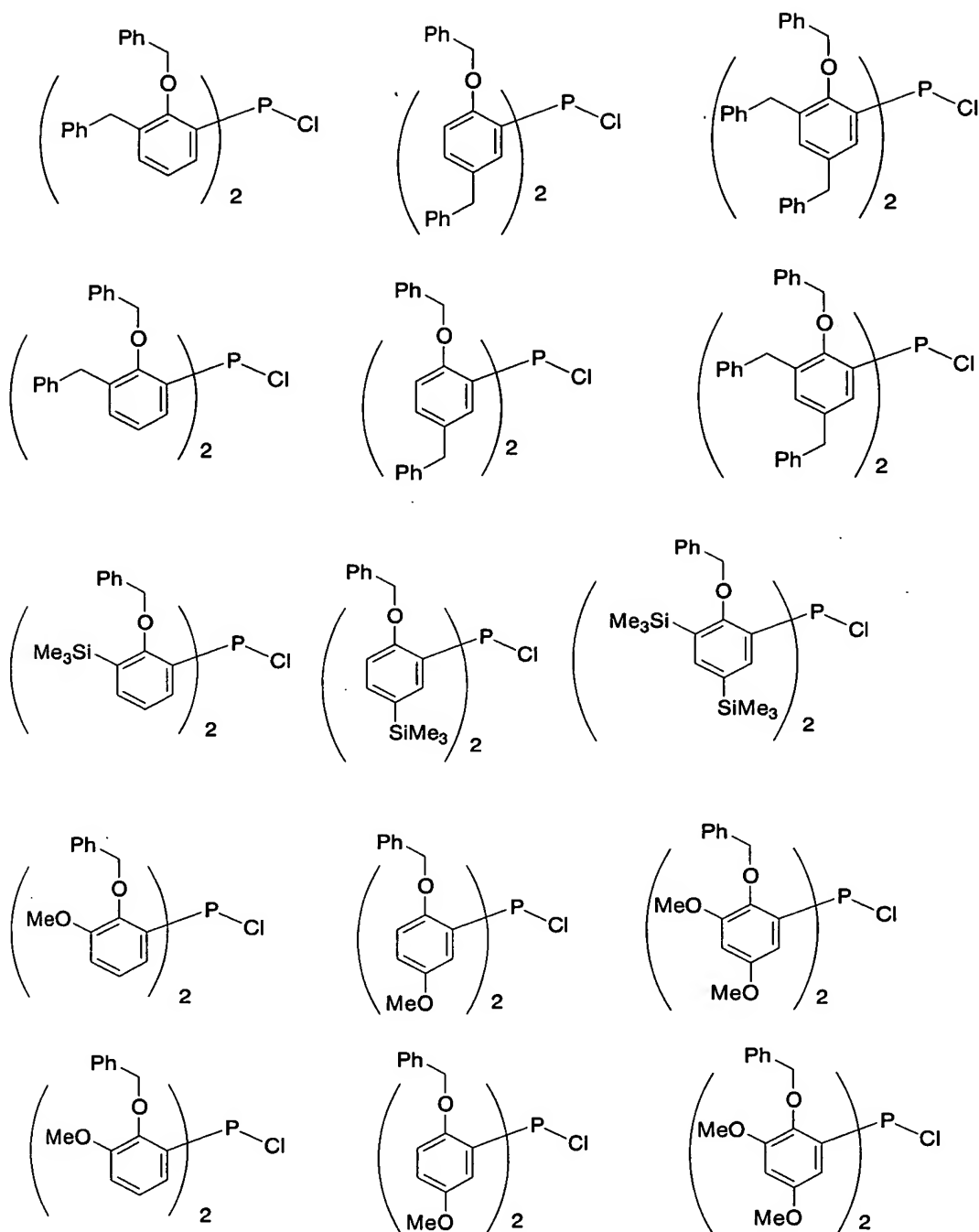
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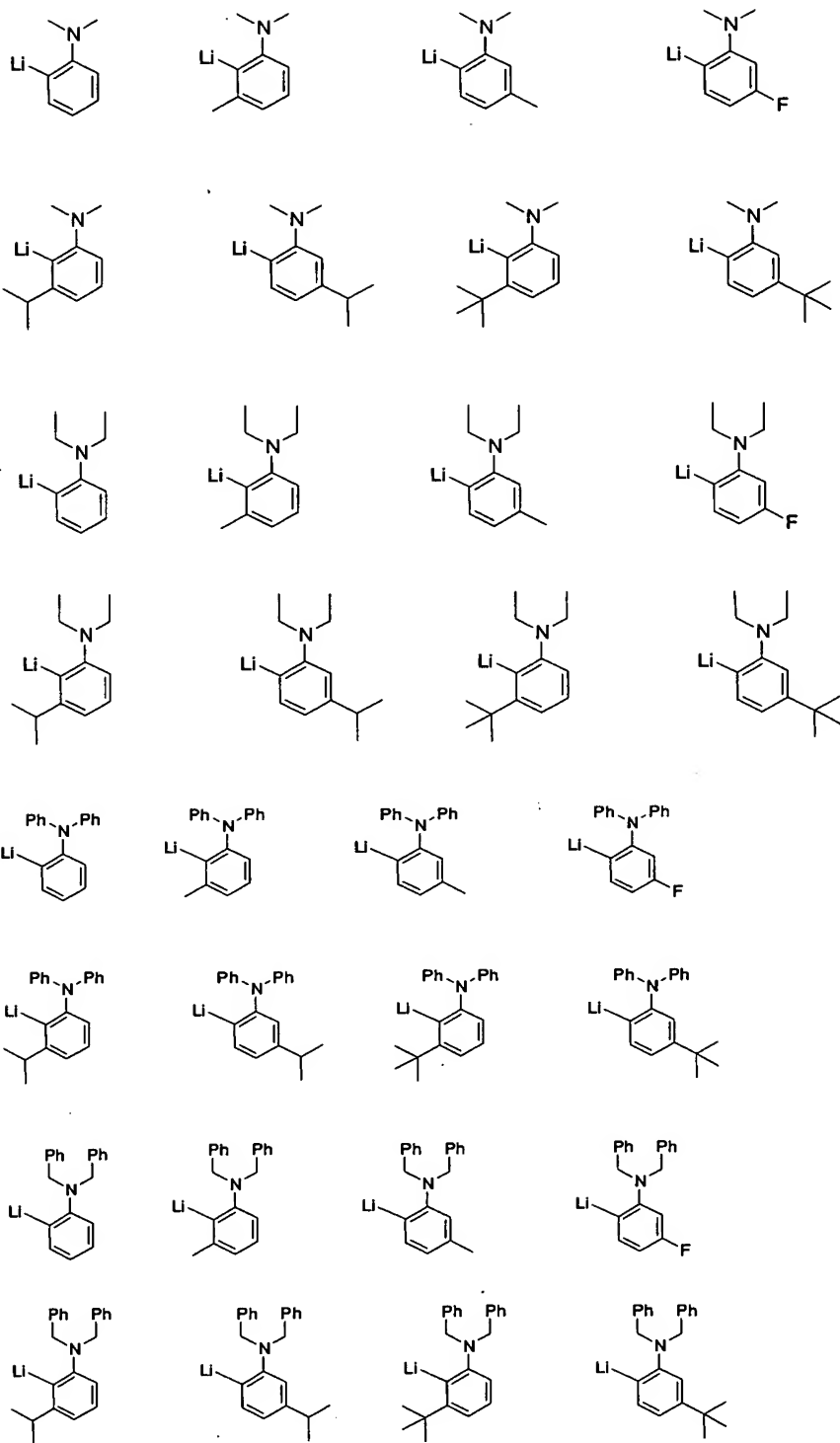


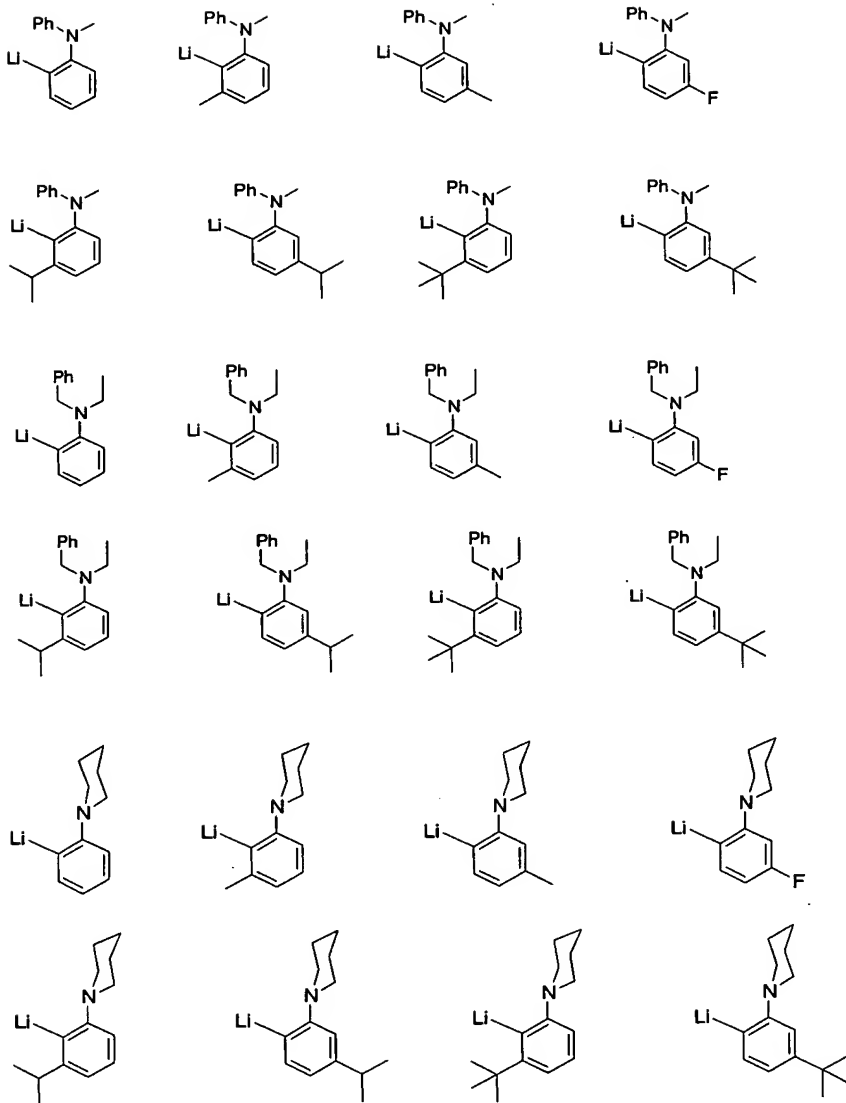


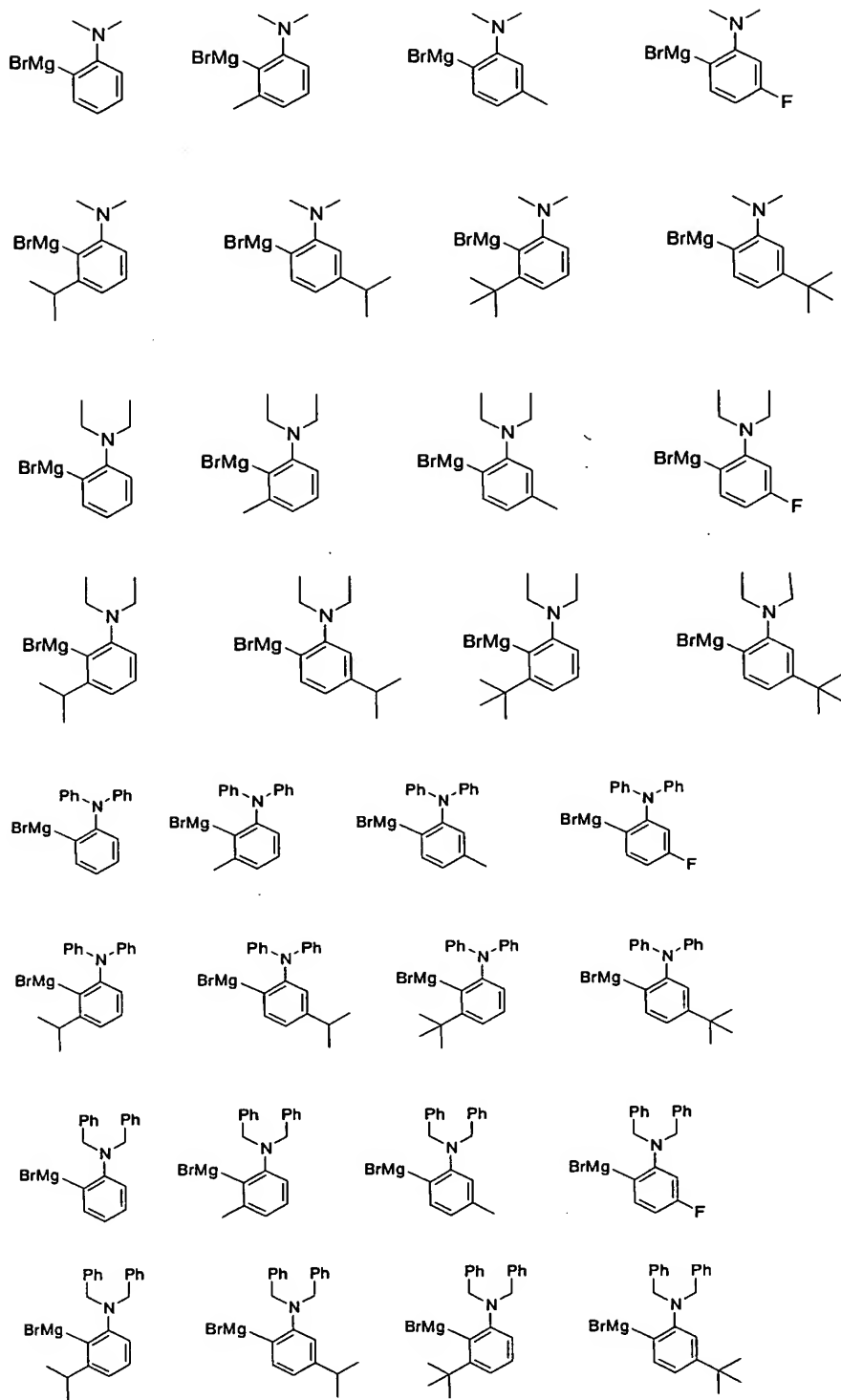
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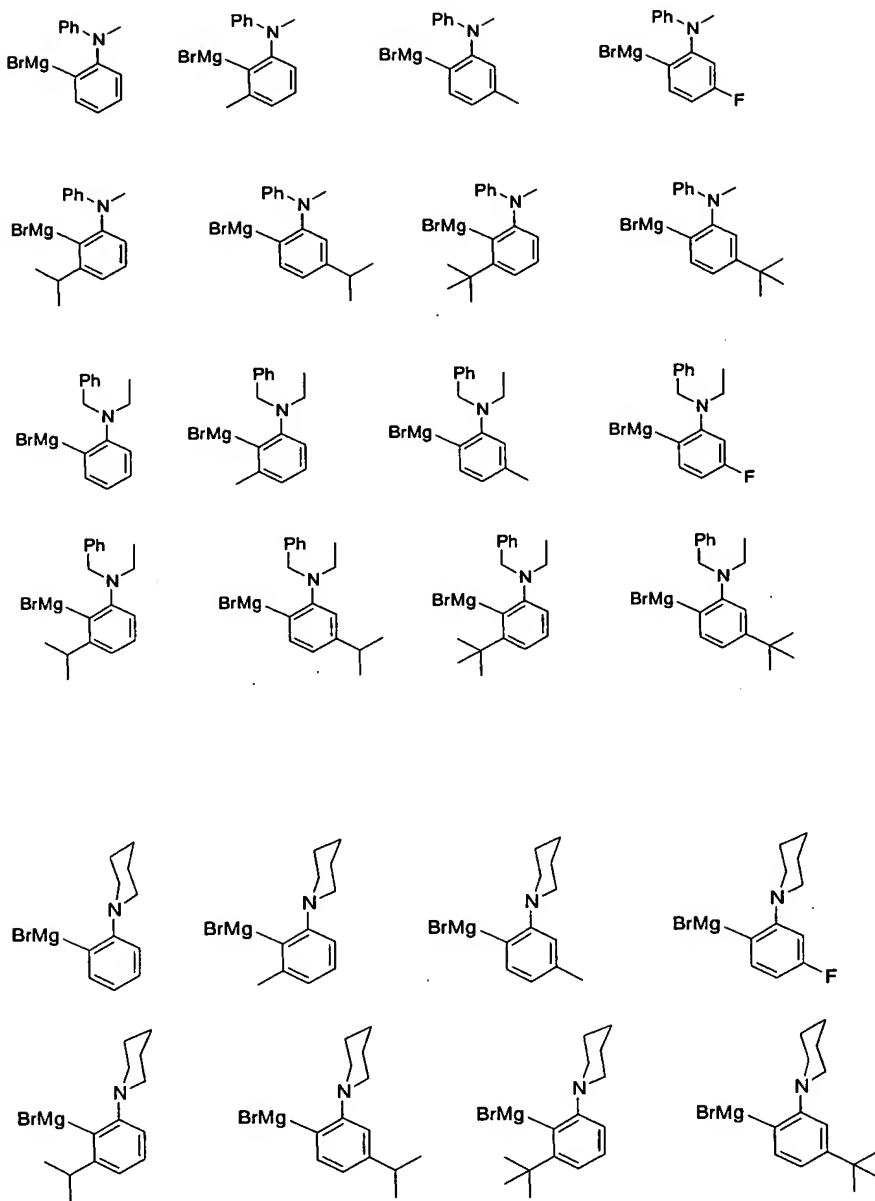


Specific examples of the aryl compound of formula (26F) include, for example, the following compounds:









Transition metal complexes of the invention obtained by
 5 reacting the ligand of formula 2 with the transition metal
 compound of formula 4 will be described below.

Examples of the group 4 element represented by M in formula
 4 include titanium, zirconium and hafnium. Titanium and zirconium
 are preferable.

Examples of the halogen atom represented by X^1 in formula 4 include fluorine, chlorine, bromine and iodine atoms. Chlorine atom is preferable.

Specific examples of the alkyl group having 1 to 10 carbon atom(s) that may be substituted, represented by X^1 , include methyl group, ethyl group, n-propyl group, isopropyl group, n-butyl group, sec-butyl group, tert-butyl group, n-pentyl group, neopentyl group, amyl group, n-hexyl group, n-octyl group and n-decyl group. Specific examples of the groups substituted with a halogen atom, alkoxy group, aryloxy group, a hydrocarbon-substituted amino group, or a silyl group substituted with a hydrocarbon include, for example, fluoromethyl group, difluoromethyl group, trifluoromethyl group, fluoroethyl group, difluoroethyl group, trifluoroethyl group, tetrafluoroethyl group, pentafluoroethyl group, perfluoropropyl group, perfluorobutyl group, perfluoropentyl group, perfluorohexyl group, perfluorooctyl perfluorodecyl trichloromethyl group, methoxymethyl group, phenoxymethyl group, dimethylaminomethyl group and trimethylsilyl group. Of the alkyl group having 1 to 10 carbon atom(s) that may be substituted, methyl, ethyl, isopropyl, tert-butyl and amyl groups are preferable, and methyl group is more preferable.

Examples of the aralkyl group having 7 to 20 carbon atoms that may be substituted, represented by X^1 , include benzyl, naphthylmethyl, anthracenylmethyl and diphenylmethyl groups, including those substituted with a halogen atom, alkyl group, alkoxy group, aryloxy group, a hydrocarbon-substituted amino group, or with a silyl group substituted with hydrocarbon. Specific examples thereof include (2-methylphenyl)methyl group,

(3-methylphenyl)methyl group, (4-methylphenyl)methyl group,
 (2,3-dimethylphenyl)methyl group, (2,4-dimethylphenyl)methyl
 group, (2,5-dimethylphenyl)methyl group,
 (2,6-dimethylphenyl)methyl group, (3,4-dimethylphenyl)methyl
 5 group, (2,3,4-trimethylphenyl)methyl group,
 (2,3,5-trimethylphenyl)methyl group,
 (2,3,6-trimethylphenyl)methyl group,
 (3,4,5-trimethylphenyl)methyl group,
 (2,4,6-trimethylphenyl)methyl group,
 10 (2,3,4,5-tetramethylphenyl)methyl group,
 (2,3,4,6-tetramethylphenyl)methyl group,
 (2,3,5,6-tetramethylphenyl)methyl group,
 (pentamethylphenyl)methyl group, (ethylphenyl)methyl group,
 (n-propylphenyl)methyl group, (isopropylphenyl)methyl group,
 15 (n-butylphenyl)methyl group, (sec-butylphenyl)methyl
 (tert-butylphenyl)methyl group, (n-pentylphenyl)methyl group,
 (neopentylphenyl)methyl group, (n-hexylphenyl)methyl
 (n-octylphenyl)methyl group, (n-decylphenyl)methyl group,
 (n-dodecylphenyl)methyl group, (fluorophenyl)methyl
 20 (difluorophenyl)methyl group, (pentafluorophenyl)methyl group,
 (chlorophenyl)methyl group, (methoxyphenyl)methyl group,
 (phenoxyphenyl)methyl group, (dimethylaminophenyl)methyl
 group and (trimethylsilylphenyl)methyl group. The aralkyl
 group having 7 to 20 carbon atoms that may be substituted is
 25 preferably a benzyl group.

Examples of the aryl group having 6 to 20 carbon atoms
 that may be substituted, represented by X^1 , include phenyl,
 naphthyl and anthracenyl groups.

Further examples of the aryl group include those

substituted a halogen atom, alkyl, alkoxy, aryloxy, a hydrocarbon-substituted amino group, or with silyl group substituted with a hydrocarbon. Specific examples thereof include 2-tolyl group, 3-tolyl group, 4-tolyl group, 2,3-xylyl group, 2,4-xylyl group, 2,5-xylyl group, 2,6-xylyl group, 3,4-xylyl group, 3,5-xylyl group, 2,3,4-trimethylphenyl group, 2,3,5-trimethylphenyl group, 2,3,6-trimethylphenyl group, 2,4,6-trimethylphenyl group, 3,4,5-trimethylphenyl group, 2,3,4,5-tetramethylphenyl group, 2,3,4,6-tetramethylphenyl group, 2,3,5,6-tetramethylphenyl group, pentamethylphenyl group, ethylphenyl group, n-propylphenyl group, isopropylphenyl group, n-butylphenyl group, sec-butylphenyl group, tert-butylphenyl group, n-pentylphenyl group, neopentylphenyl group, n-hexylphenyl group, n-octylphenyl group, n-decylphenyl group, n-dodecylphenyl group, n-tetradecylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 3,5-difluorophenyl group, pentafluorophenyl group, 4-chlorophenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 4-phenoxyphenyl group, 4-dimethylaminophenol group and 4-trimethylsilylphenyl group. Aryl group that may be substituted is preferably a phenyl group.

Specific examples of the alkoxy group having 1 to 10 carbon atom(s) that may be substituted, represented by X^1 , include methoxy group, ethoxy group, n-propoxy group, isopropoxy group, n-butoxy group, sec-butoxy group, tert-butoxy group, n-pentyloxy group, neopentyloxy group, n-hexyloxy group, n-octyloxy group, n-nonyloxy group and n-decyloxy group. These groups may be further substituted, and examples thereof include

those substituted with a halogen atom, an alkoxy, aryloxy, hydrocarbon-substituted amino, or with a silyl group substituted with a hydrocarbon.

Specific examples of the substituted alkoxy group include
5 fluoromethoxy group, fluoroethoxy group, difluoromethoxy group, trifluoromethoxy group, fluoroethoxy group, difluoroethoxy group, trifluoroethoxy group, tetrafluoroethoxy group, pentafluoroethoxy group, perfluoropropoxy group, perfluorobutyloxy group, perfluoropentyloxy group,
10 perfluorohexyloxy group, perfluorooctyloxy group, perfluorodecyloxy group, trichloromethyloxy group, methoxymethoxy group, phenoxymethoxy group, dimethylamiomethoxy group, trimethylsilylmethoxy group and the like. The alkoxy group having 1 to 10 carbon atom(s) that may
15 be substituted is preferably a methoxy group.

Examples of the aralkyloxy group having 7 to 20 carbon atoms, represented by X^1 , include benzyloxy group, naphthylmethoxy group, anthracenylmethoxy group and diphenylmethoxy group. These may be further substituted, and
20 examples thereof include those substituted with a halogen atom, an alkyl group, an alkoxy group, an aryloxy group, a hydrocarbon-substituted amino group, or with a silyl' group substituted with a hydrocarbon. Specific examples thereof include (2-methylphenyl)methoxy group,
25 (3-methylphenyl)methoxy group, (4-methylphenyl)methoxy group, (2,3-dimethylphenyl)methoxy group, (2,4-dimethylphenyl)methoxy group, (2,5-dimethylphenyl)methoxy group, (2,6-dimethylphenyl)methoxy group,

(3,4-dimethylphenyl)methoxy group,
 (2,3,4-trimethylphenyl)methoxy group,
 (2,3,5-trimethylphenyl)methoxy group,
 (2,3,6-trimethylphenyl)methoxy group,
 5 (3,4,5-trimethylphenyl)methoxy group,
 (2,4,6-trimethylphenyl)methoxy
 group, (2,3,4,5-tetramethylphenyl)methoxy group,
 (2,3,4,6-tetramethylphenyl)methoxy group,
 (2,3,5,6-tetramethylphenyl)methoxy group,
 10 (pentamethylphenyl)methoxy group, (ethylphenyl)methoxy group,
 (n-propylphenyl)methoxy group, (isopropylphenyl)methoxy group,
 (n-butylphenyl)methoxy group, (sec-butylphenyl)methoxy group,
 (tert-butylphenyl)methoxy group, (n-pentylphenyl)methoxy
 group, (neopentylphenyl)methoxy group, (n-hexylphenyl)methoxy
 15 group, (n-octylphenyl)methoxy group, (n-decylphenyl)methoxy
 group, (n-dodecylphenyl)methoxy group, (fluorophenyl)methyl
 group, (difluorophenyl)methyl group,
 (pentafluorophenyl)methyl group, (chlorophenyl)methyl group,
 (methoxyphenyl)methyl group, (phenoxyphenyl)methyl
 20 (dimethoxyaminophenyl)methyl group and
 (trimethoxysilylphenyl)methyl group. The aralkyloxy group
 havin 7 to 20 carbon atoms that may be substituted is preferably
 a benzyloxy group.

Examples of the aryloxy group having 6 to 20 carbon atoms
 25 that may be substituted in X¹ include phenoxy, naphthoxy and
 anthracenoxy groups. These groups may be further substituted,
 and examples thereof include those substituted with a halogen
 atom, an alkyl group, an alkoxy group, an aryloxy group, a
 hydrocarbon-substituted amino group, or with a silyl group

substituted with a hydrocarbon. Specific examples thereof include 2-methylphenoxy group, 3-methylphenoxy group, 4-methylphenoxy group, 2,3-dimethylphenoxy group, 2,4-dimethylphenoxy group, 2,5-dimethylphenoxy group, 2,6-dimethylphenoxy group, 3,4-dimethylphenoxy group, 3,5-dimethylphenoxy group, 2,3,4-trimethylphenoxy group, 2,3,5-trimethylphenoxy group, 2,3,6-trimethylphenoxy group, 2,4,5-trimethylphenoxy group, 2,4,6-trimethylphenoxy group, 3,4,5-trimethylphenoxy group, 2,3,4,5-tetramethylphenoxy group, 2,3,4,6-tetramethylphenoxy group, 2,3,5,6-tetramethylphenoxy group, pentamethylphenoxy group, ethylphenoxy group, n-propylphenoxy group, isopropylphenoxy group, n-butylphenoxy group, sec-butylphenoxy group, tert-butylphenoxy group, n-hexylphenoxy group, n-octylphenoxy group, n-decylphenoxy group, n-tetradecylphenoxy group, 2-fluorophenoxy group, 3-fluorophenoxy group, 4-fluorophenoxy group, 3,5-difluorophenoxy group, pentafluorophenoxy group, 4-chlorophenoxy group, 2-methoxyphenoxy group, 3-methoxyphenoxy group, 4-methoxyphenoxy group, 4-phenoxyphenoxy group, 4-dimethylaminophenoxy group and 4-trimethylsilylphenoxy group. The aryloxy group having 7 to 20 carbon atoms that may be substituted is preferably a phenoxy group.

In the amino group disubstituted with substituted or unsubstituted hydrocarbon group having 1 to 20 carbon atom(s), represented by X^1 , examples of the hydrocarbon include alkyl groups having 1 to 10 carbon atom(s) such as methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, tert-butyl, n-pentyl, neopentyl, amyl, n-hexyl, cyclohexyl, n-octyl, or n-decyl group;

and aryl groups having 6 to 20 carbon atoms such as phenyl, tolyl, xylyl, naphthyl or anthracenyl group. Examples of the amino group substituted with the hydrocarbon groups having 1 to 20 carbon atom(s) include dimethylamino group, diethylamino group, 5 di-n-propylamino group, diisopropylamino group, di-n-butylamino group, di-sec-butylamino group, di-tert-butylamino group, di-isobutylamino group, tert-butylisopropylamino group, di-n-hexylamino group, di-n-octylamino group, di-n-decylamino group, diphenylamine 10 group and the like. The dimethylamino and diethylamino groups are preferable.

The neutral ligand represented by L or L¹ denotes a molecule having a neutral functional group(s) such as ether, sulfide, amine, phosphine or olefin, and it may possess coordinating 15 functional groups at a plurality of sites in the molecule.

Examples of the neutral ligand include dimethyl ether, diethyl ether, methyl tert-butyl ether, furan, tetrahydrofuran, dimethoxyethane, diethoxyethane, dimethyl sulfide, diethyl sulfide, methyl tert-butyl sulfide, thiophene, 20 tertahydrothiophene, ethylenedithiol, dimethylsulfide, ethylenedithiol, diethylsulfide, trimethylamine, triethylamine, triphenylamine, tricyclohexylamine, pyridine, 2,2'-bipyridine, tetramethylenediamine, tetraethylethylenediamine, triphenylphosphine, 25 tricyclohexylphosphine, tri-tert-butylphosphine, bis(diphenylphosphino)methane, bis(diphenylphosphino)ethane, bis(diphneylphosphino)propane, bis(diphneylphosphino)binaphthyl, ethylene, propylene, butene, butadiene, octene, octadiene, cyclohexene, cyclohexadiene,

norbornene and norbornadiene.

Examples of the transition metal compound of formula (4) include, for example, tetrabenzyl titanium, tetraneopentyl titanium, tetrachloro titanium, tetraisopropoxy titanium, 5 diisopropoxy titanium dichloride, tetrakis(dimethylamino)titanium, tetrakis(diethylamino)titanium, bis(dimethylamino)titanium dichloride, bis(diethylamino)titanium dichloride, tetrakis(trifluoroacetoxy)titanium, 10 bis(trifluoroacetoxy)titanium dichloride, titanium trichloride-3tetrahydrofuran complex, titanium tetrachloride-2tetrahydrofuran complex, tetrabenzyl zirconium, tetraneopentyl zirconium, tetrachlorozirconium, tetraisopropoxy zirconium, diisopropoxy 15 zirconium dichloride, tetrakis(dimethylamino) zirconium, tetrakis(diethylamino) zirconium, bis(dimethylamino) zirconium dichloride, bis(diethylamino)zirconium dichloride, tetrakis(trifluoroacetoxy) zirconium, bis(trifluoroacetoxy) zirconium dichloride, trichlorozirconium-3tetrahydrofuran 20 complex, tetrachlorozirconium-2tetrahydrofuran complex, tetrabenzyl hafnium, tetraneopentyl hafnium, tetrachlorohafnium, tetraisopropoxy hafnium, diisopropoxy hafnium dichloride, tetrakis(dimethylamino)hafnium, tetrakis(diethylamino)hafnium, bis(dimethylamino)hafnium 25 dichloride, bis(diethylamino)hafnium dichloride, tetrakis(trifluoroacetoxy)hafnium, bis(trifluoroacetoxy)hafnium, trichlorohafnium 3-tetrahydrofuran complex, tetrachlorohafnium 2-tetrahydrofuran complex and the like.

The transition metal complex is produced by reacting phosphine compound of formula 2 with the transition metal compound of formula 4.

5 The ratio between the phosphine compound of formula (2) and the transition metal compound of formula (4) is not particularly restricted, and it is preferably in the range of 1:0.1 to 1:10, more preferably in the range of 1:0.5 to 1:2.

A base is used for the reaction, if necessary. Examples of the base include organic alkali metal compounds including
10 organic lithium compounds such as methyl lithium, ethyl lithium, n-butyl lithium, sec-butyl lithium, tert-butyl lithium, lithium trimethylsilylacetylide, lithium acetylide, trimethylsilylmethyl lithium, vinyl lithium, phenyl lithium, allyl lithium or the like, and metal hydrides such as sodium
15 hydride, potassium hydride or the like. The amount thereof is usually in the range of 0.5 to 5 mole per mol of the phosphine compound of formula (2).

The reaction is usually carried out in a solvent inert to the reaction. Examples of the solvent include aprotic
20 solvents including aromatic hydrocarbon solvents such as benzene, toluene or the like; aliphatic hydrocarbon solvents such as hexane, heptane or the like; and ether solvents such as diethyl ether, tetrahydrofuran, 1,4-dioxane or the like; amide solvents such as hexamethylphosphoric amide, dimethylformamide or the
25 like; polar solvents such as acetonitrile, propionitrile, acetone, diethyl ketone, methyl isobutyl ketone, cyclohexanone or the like; and halogenated solvents such as dichloromethane, dichloroethane, chlorobenzene, dichlorobenzene or the like. These solvents may be used alone or as a mixture of at least

two of them. The amount thereof is usually 1 to 200 parts by weight, preferably 3 to 50 parts by weight, per part by weight of the phosphine compound of formula (2).

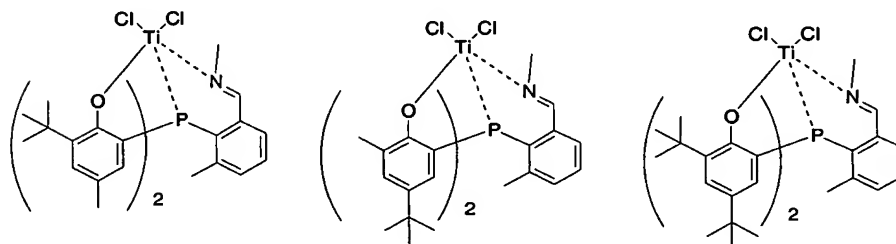
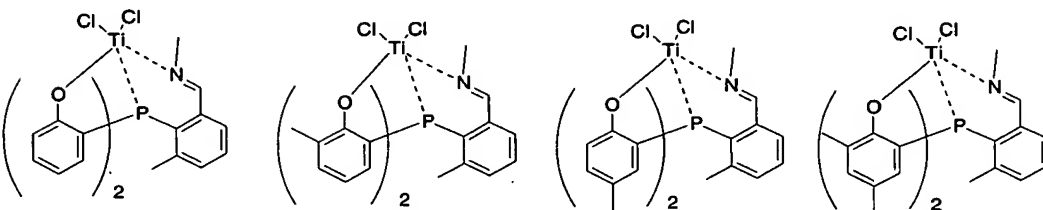
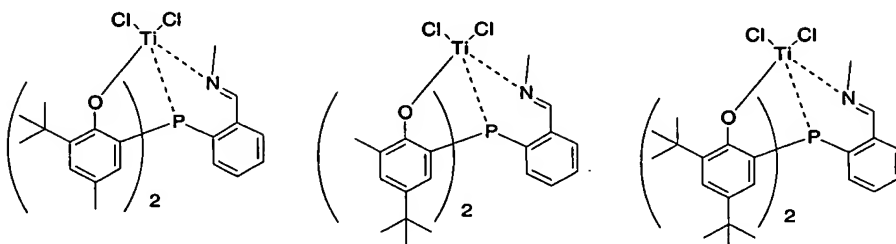
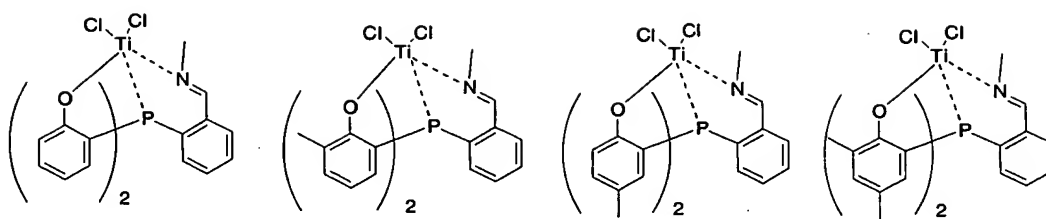
The reaction is usually carried out by adding the transition metal compound of formula (4) to the phosphine compound of formula (2) in a solvent after adding a base, if necessary. The reaction temperature is usually in the range of -100°C or more to the boiling point or less of the solvent, preferably in the range of -80 to 100°C.

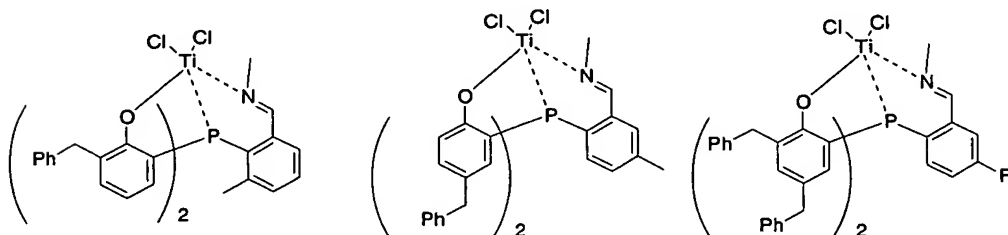
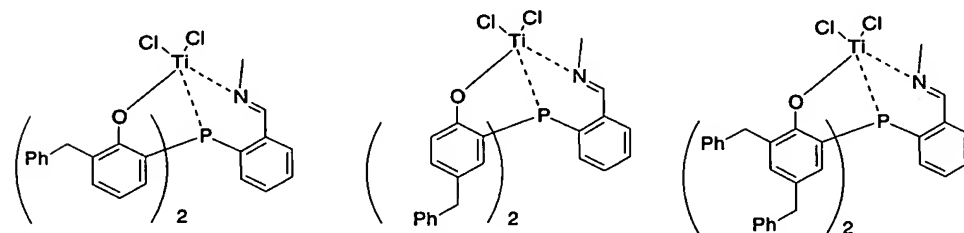
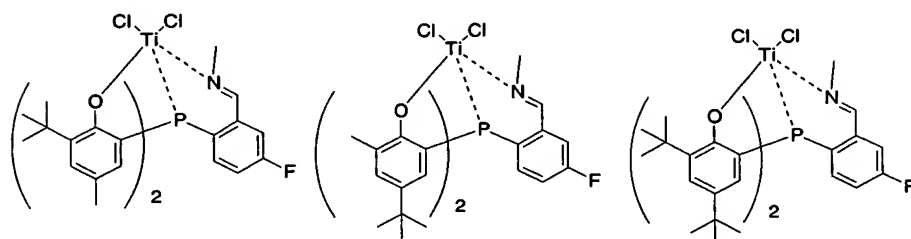
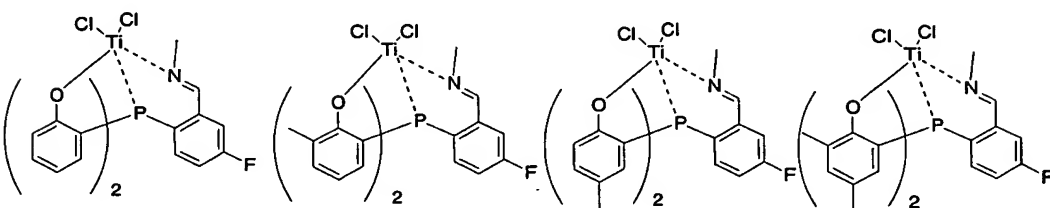
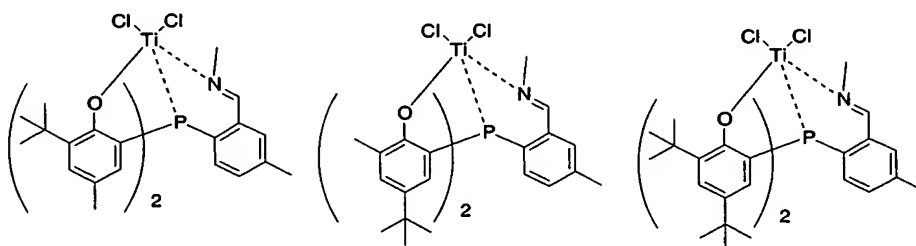
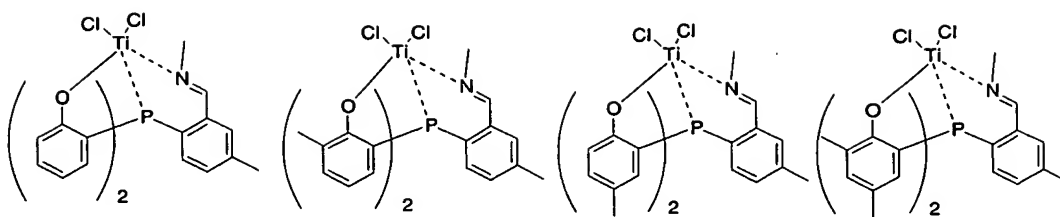
The transition metal compound may be obtained from the reaction mixture by a conventional method. For example, the precipitated substance is removed by filtration, and a solid product is precipitated by concentrating the filtrate.

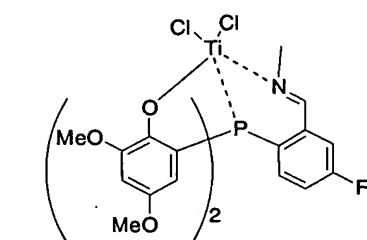
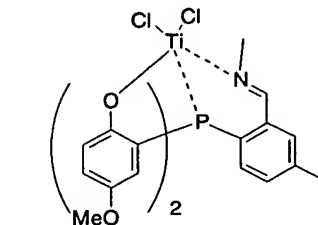
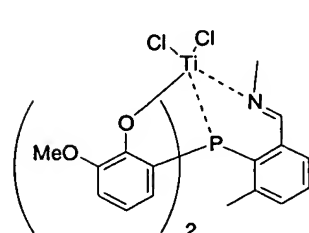
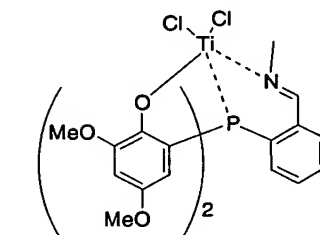
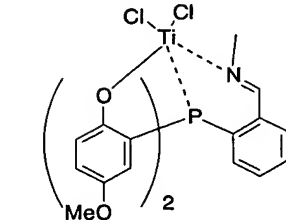
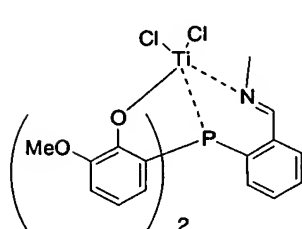
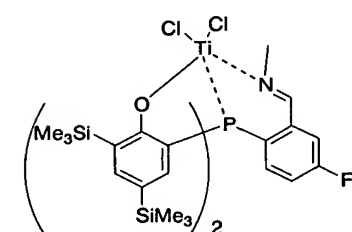
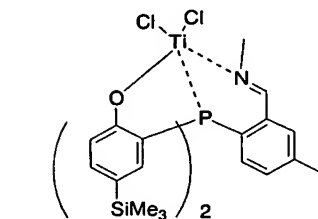
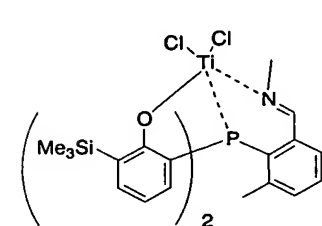
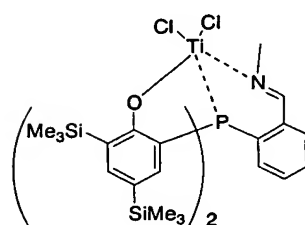
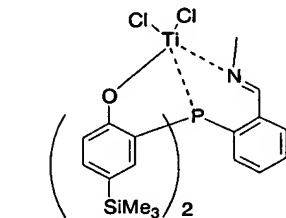
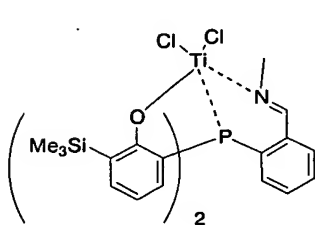
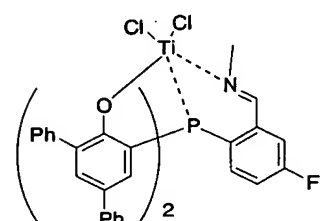
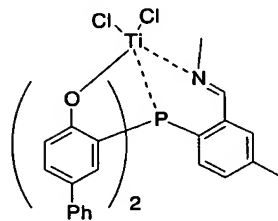
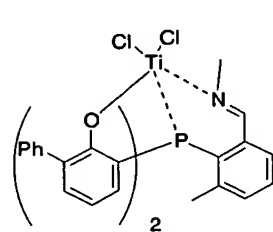
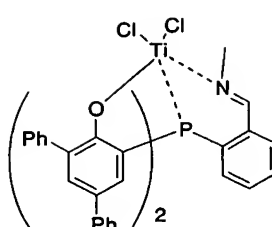
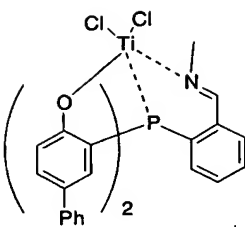
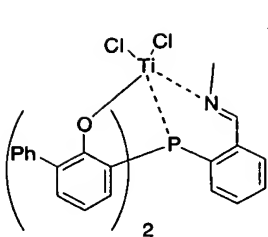
The transition metal compound thus obtained is typically the transition metal compound of formula (3).

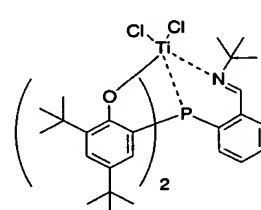
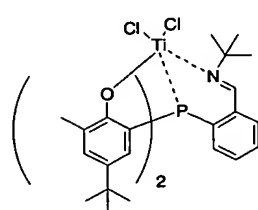
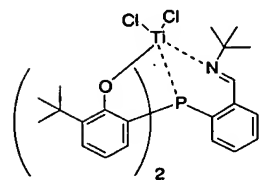
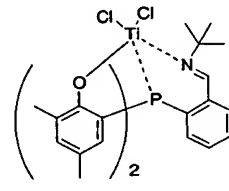
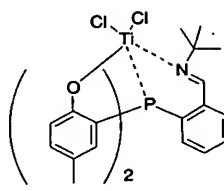
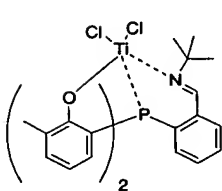
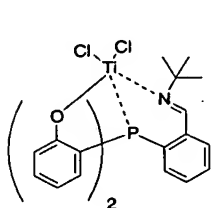
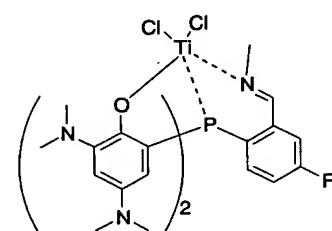
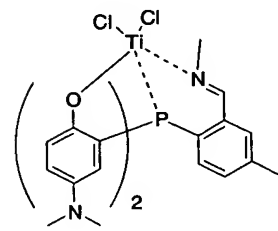
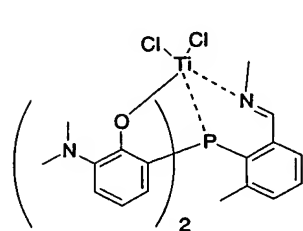
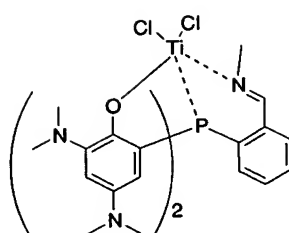
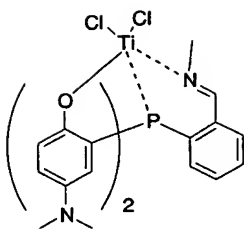
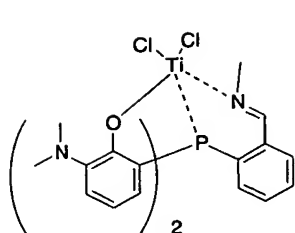
Examples of the compound having a partial structure corresponding to each formula of G^2 in formula (3) include the following compounds.

Examples of the transition metal compound of formula (3) include, for example, the following compounds:

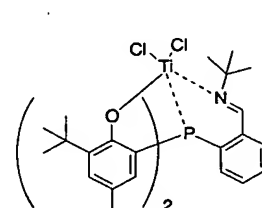
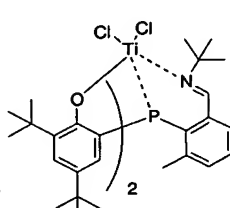
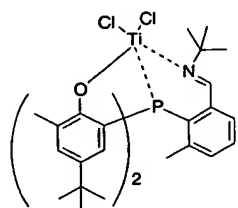
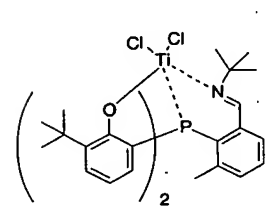
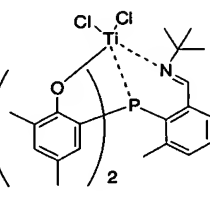
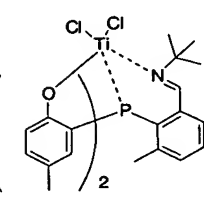
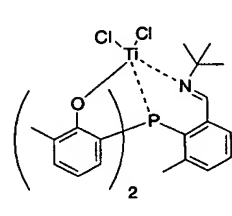
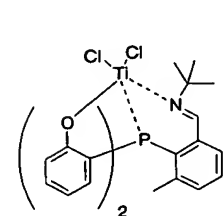


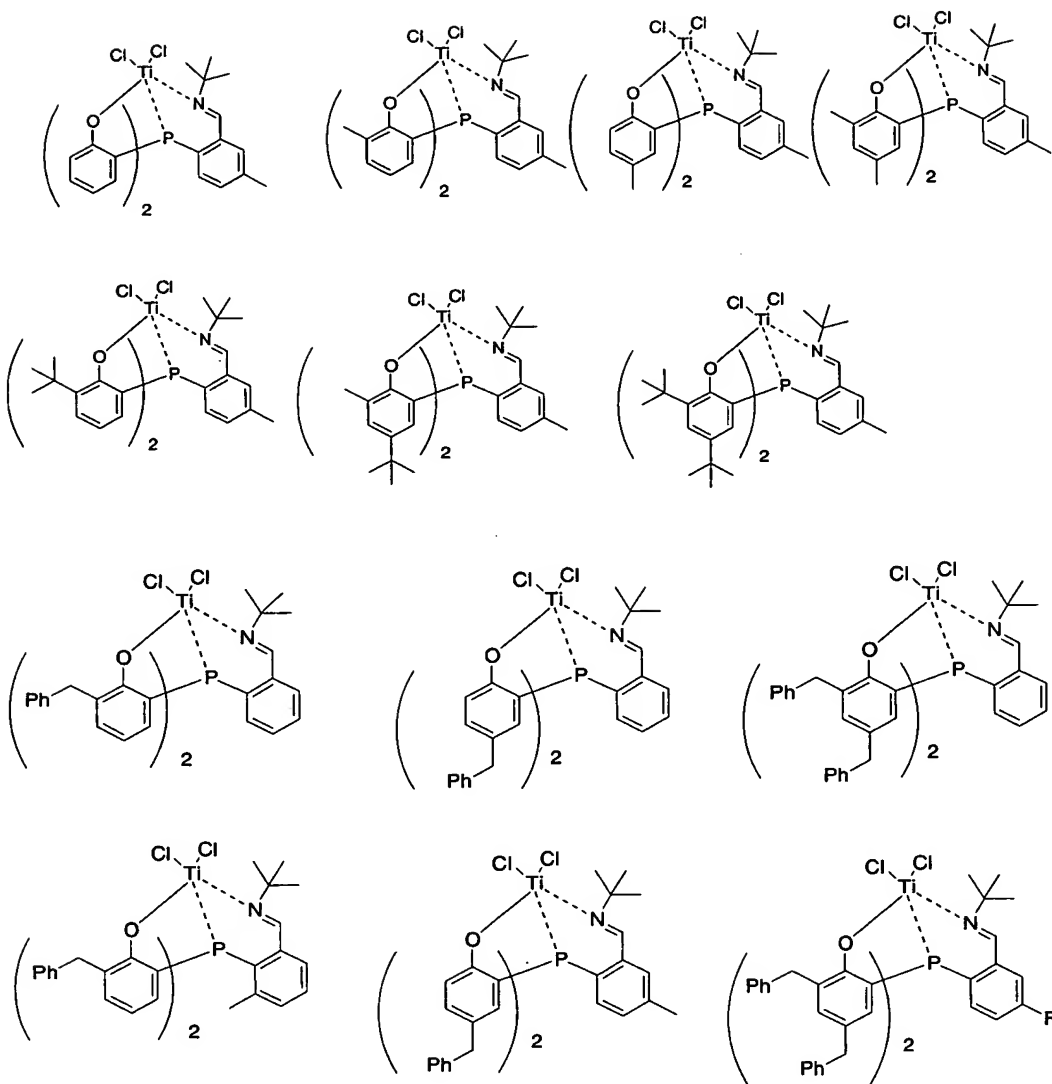


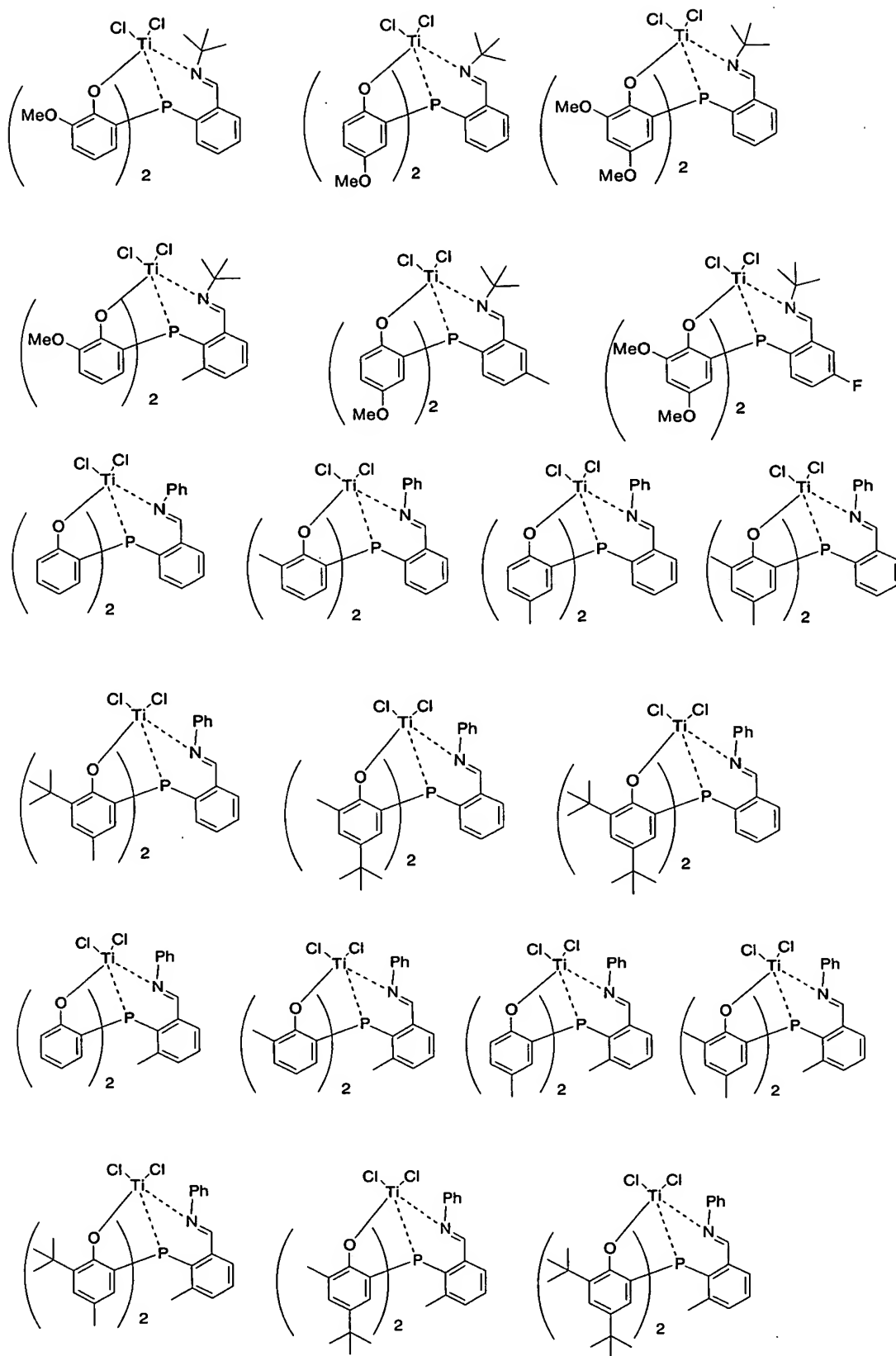


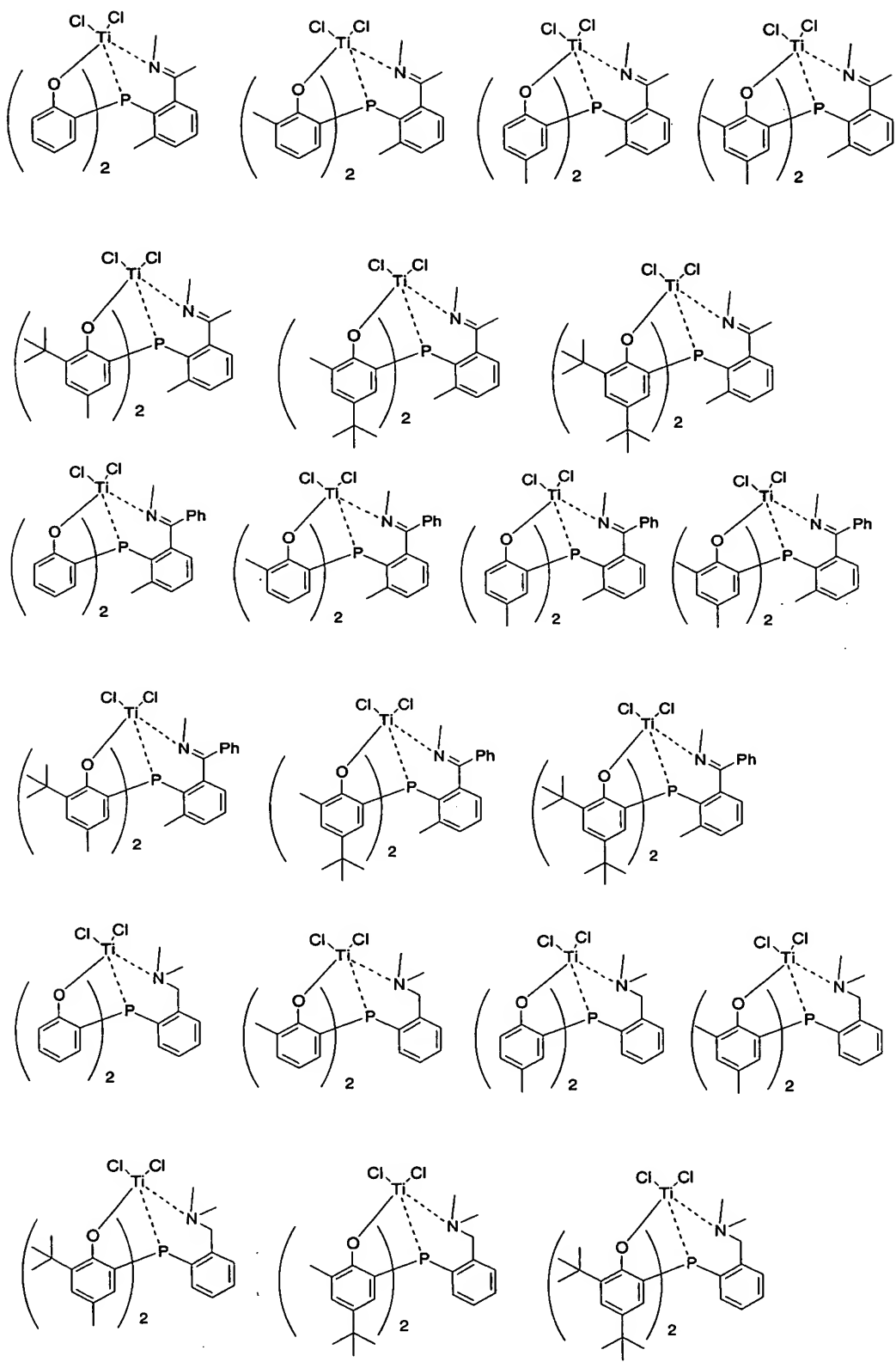


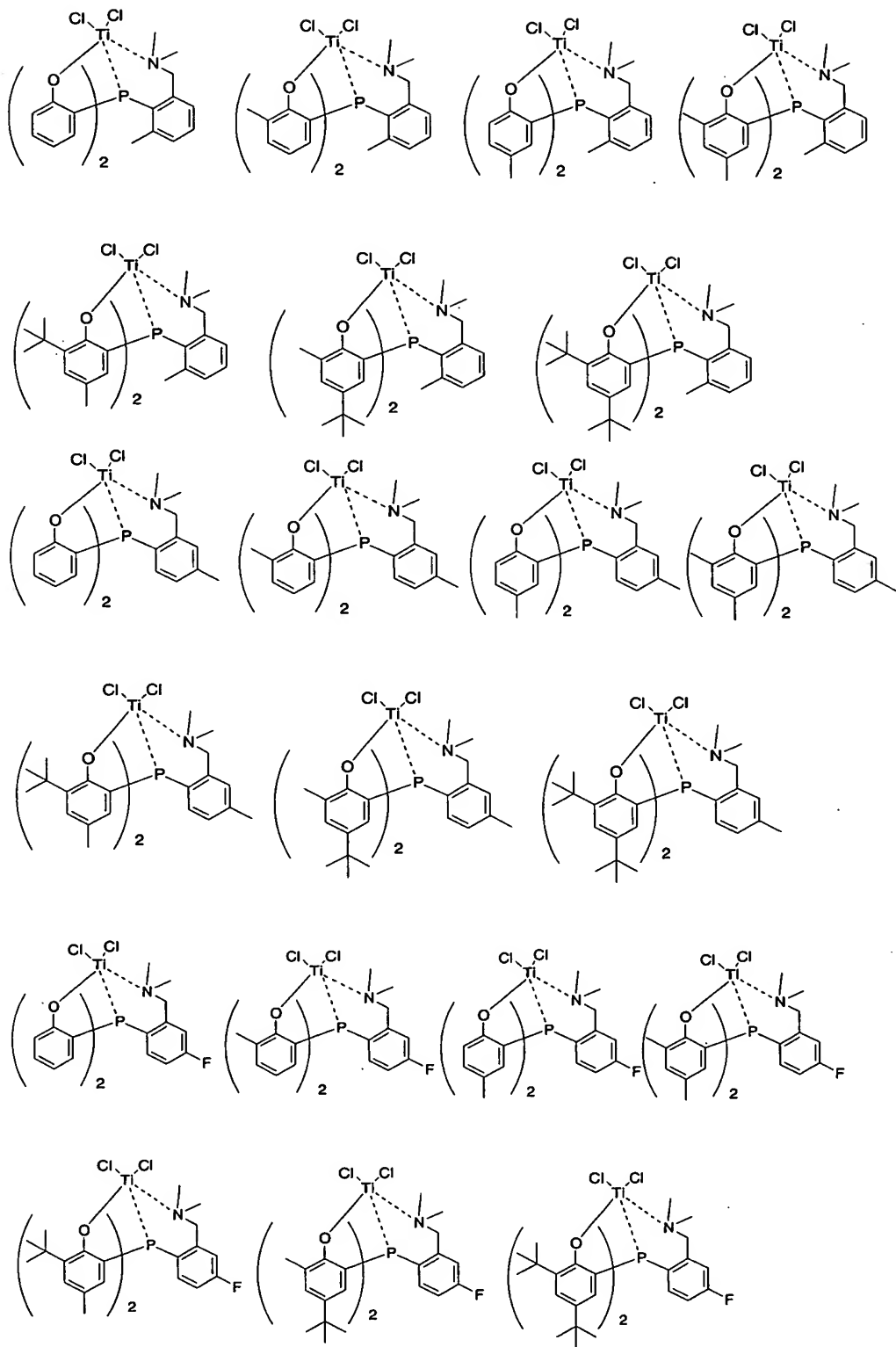
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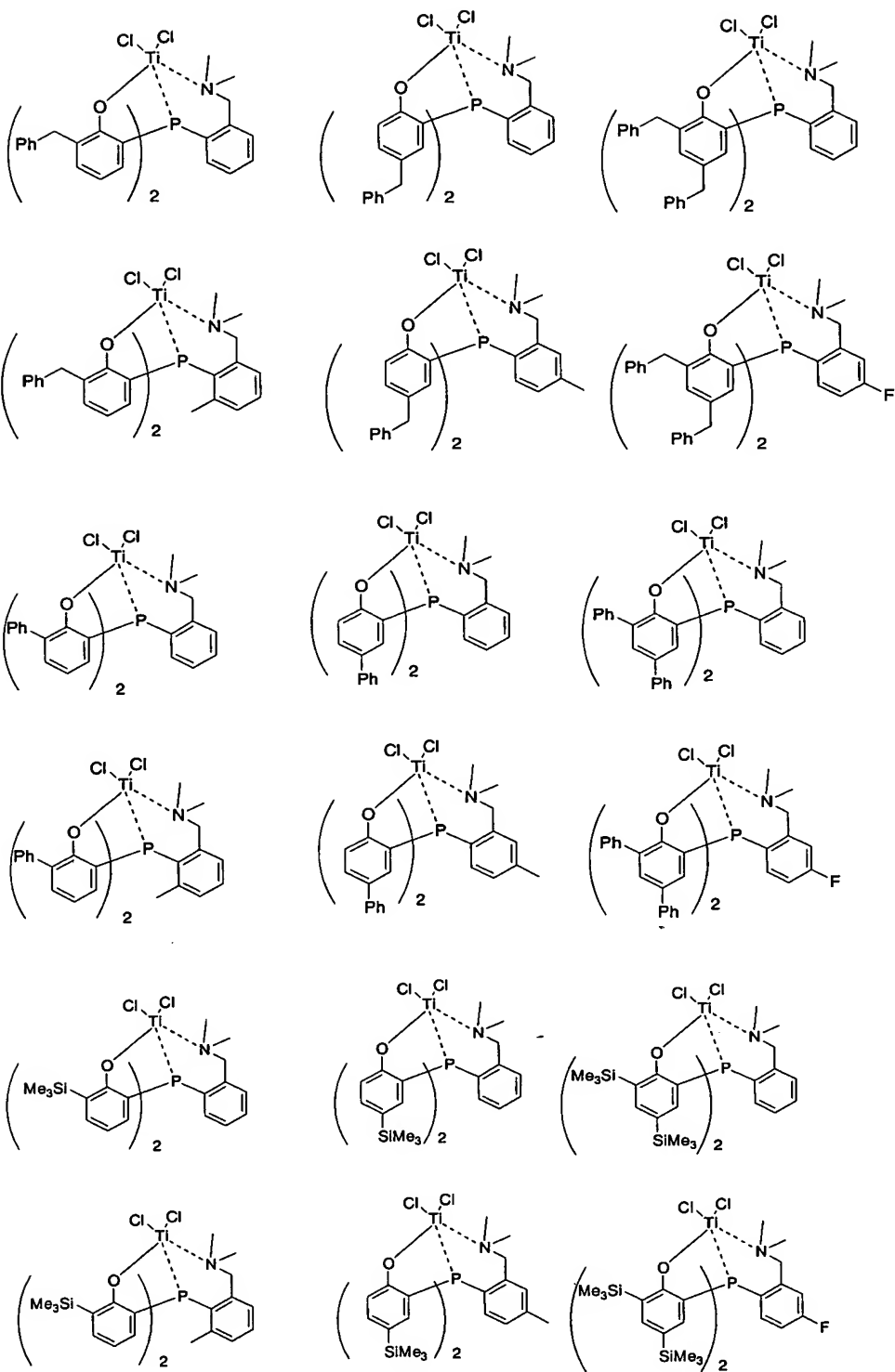


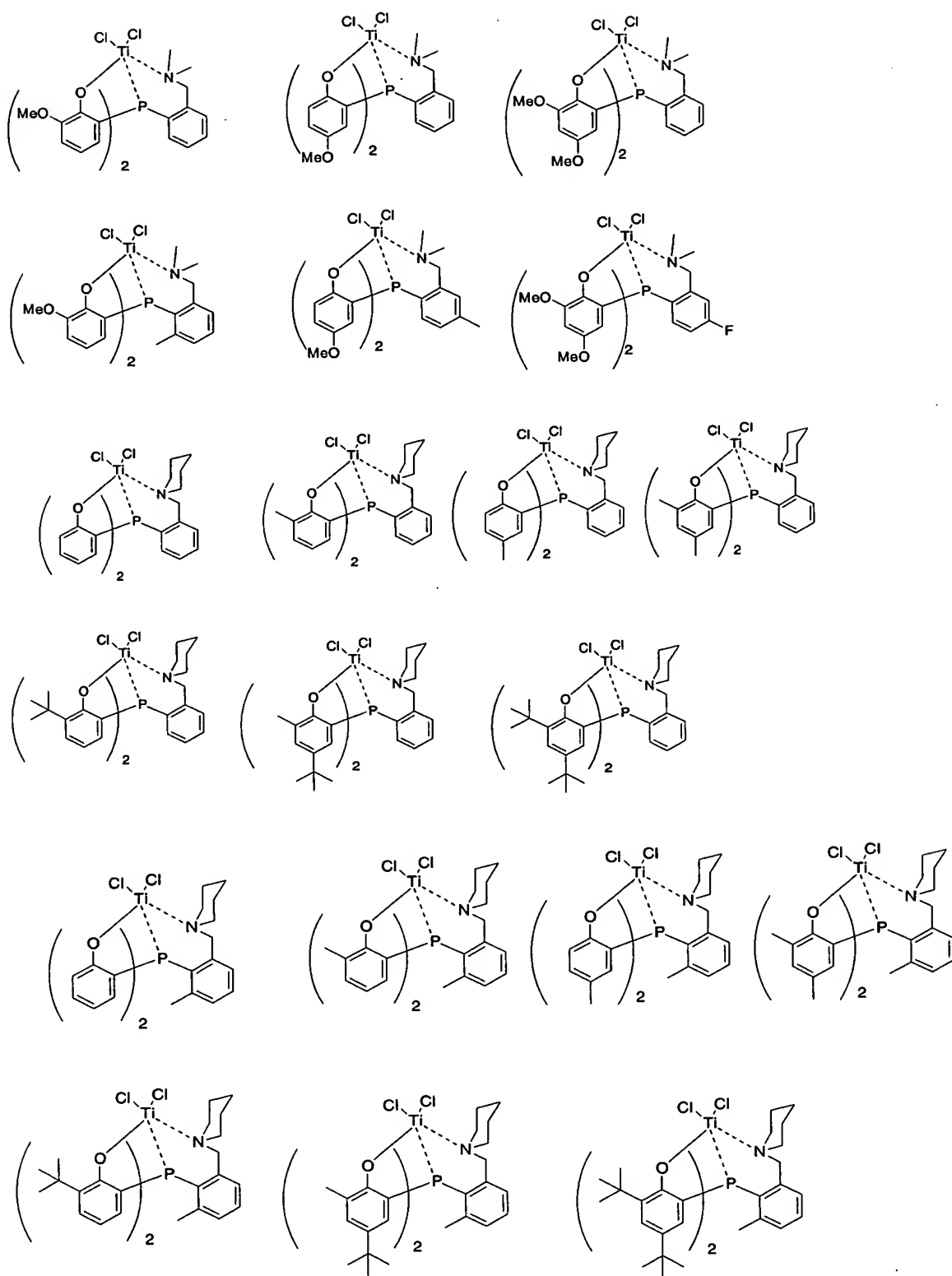


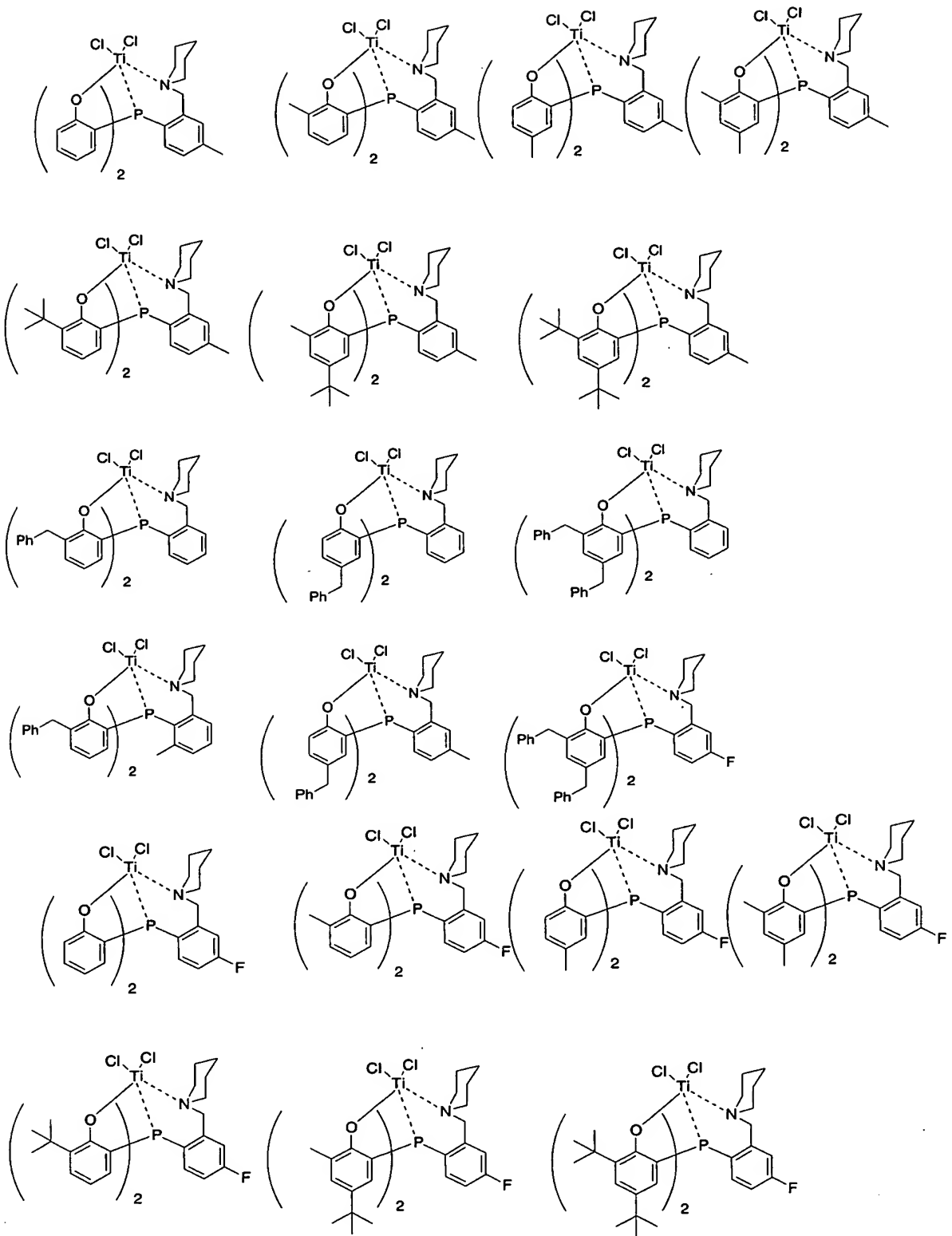


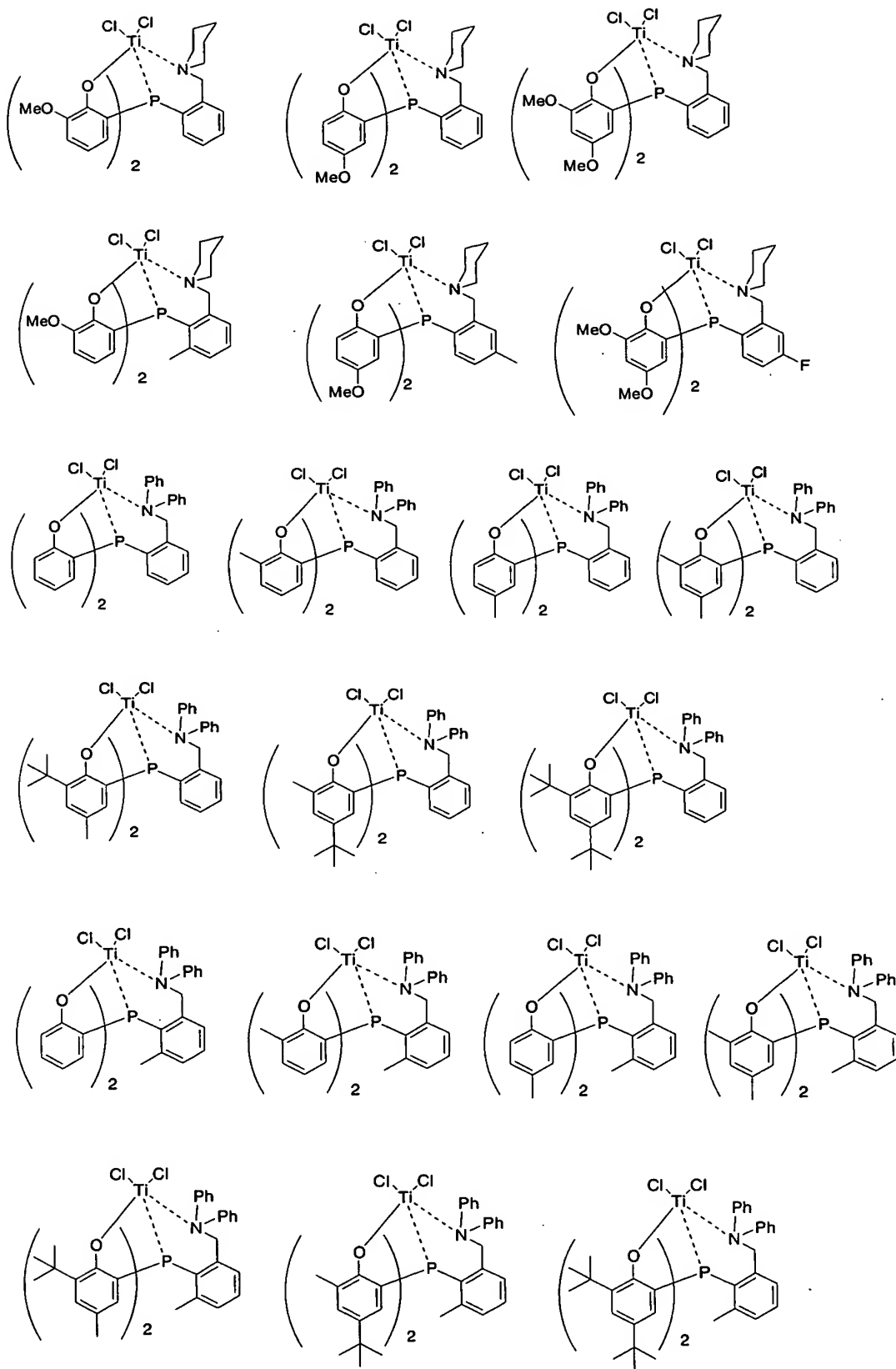


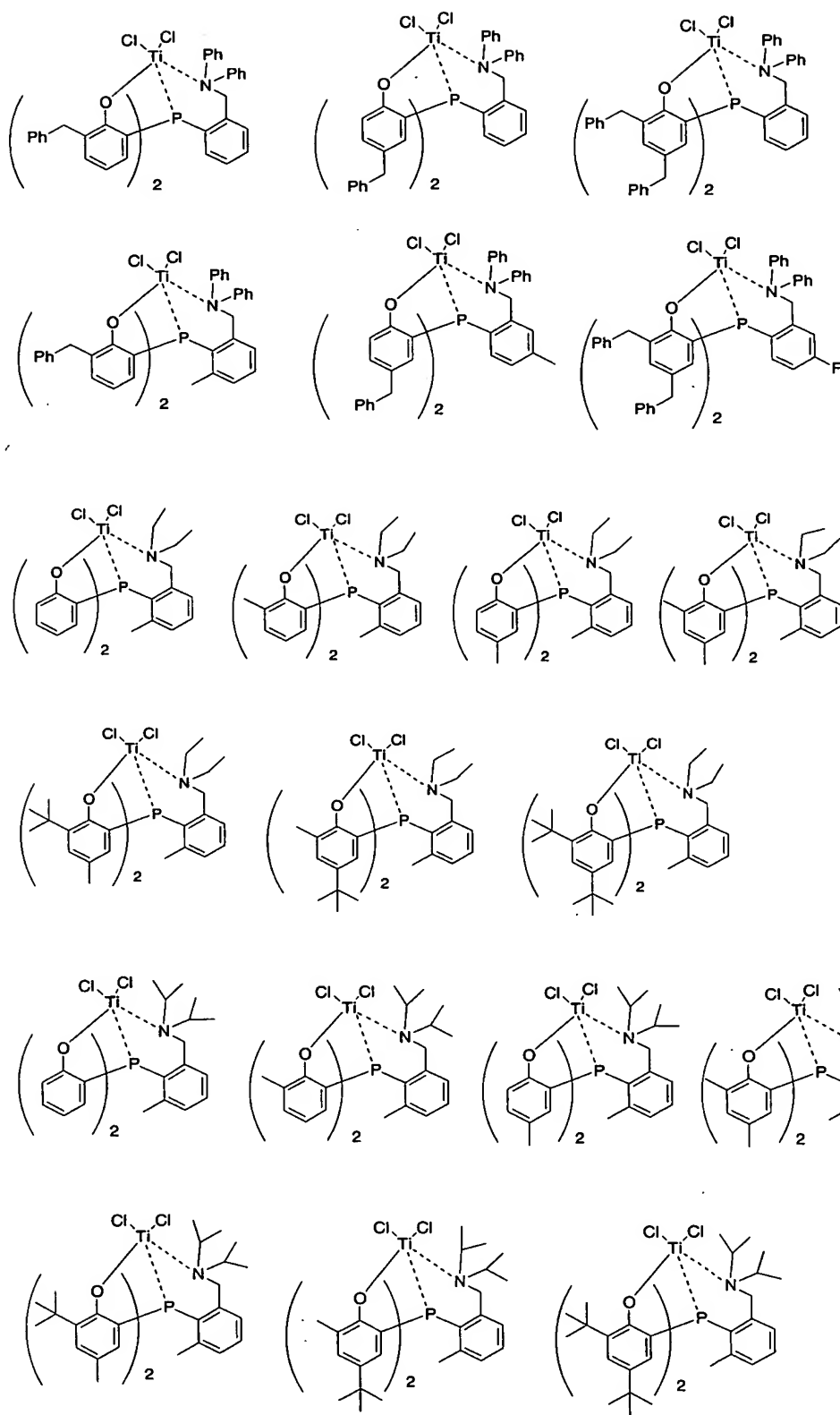


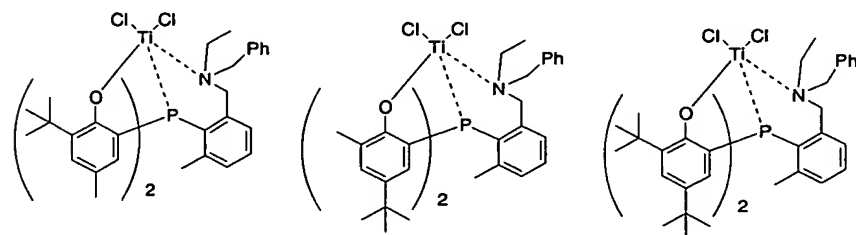
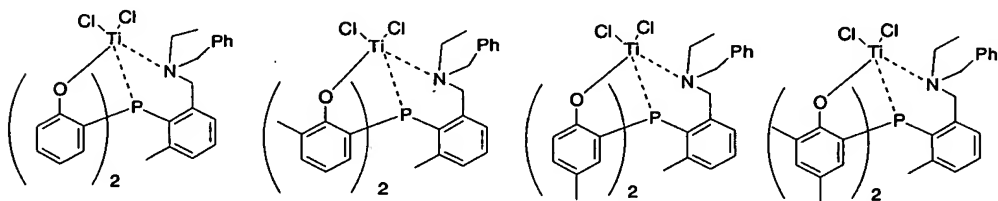
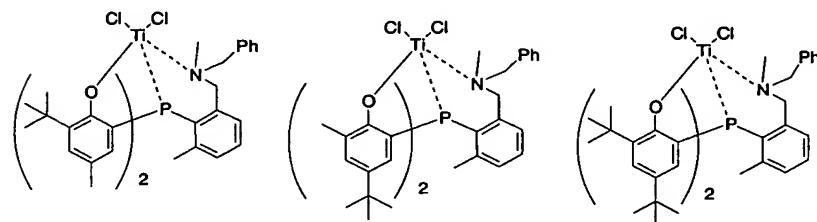
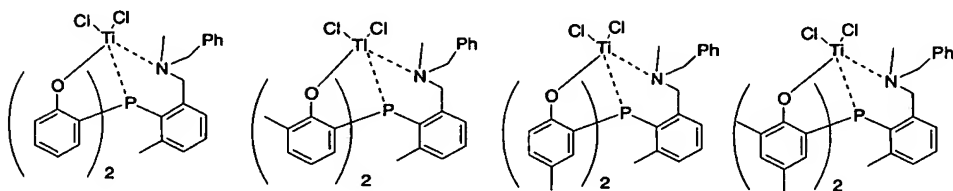
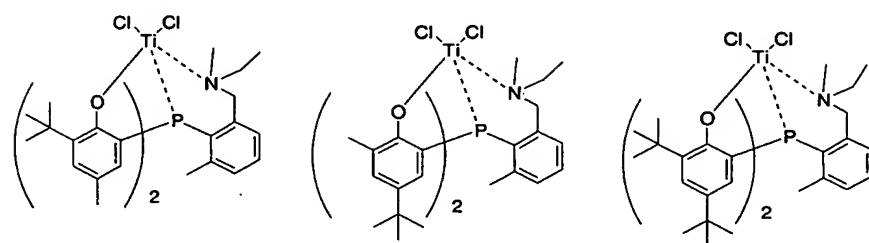
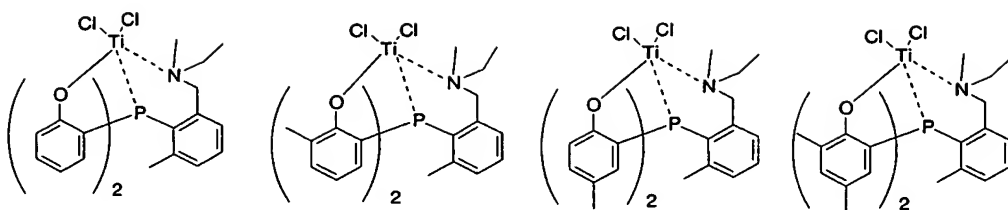


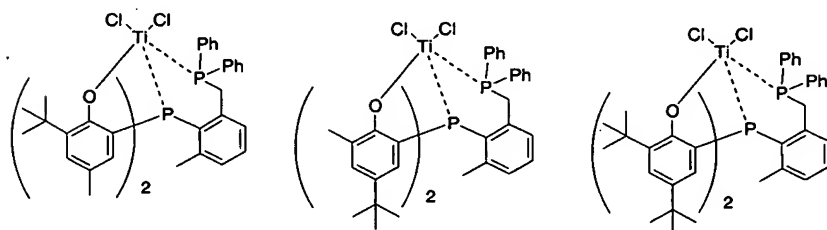
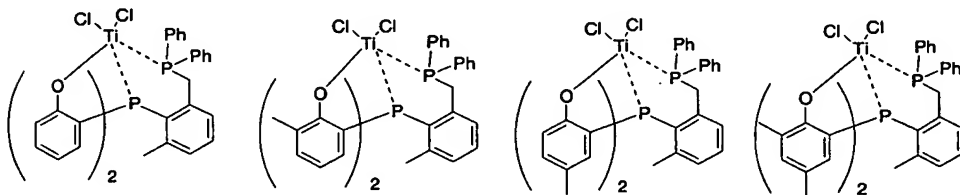
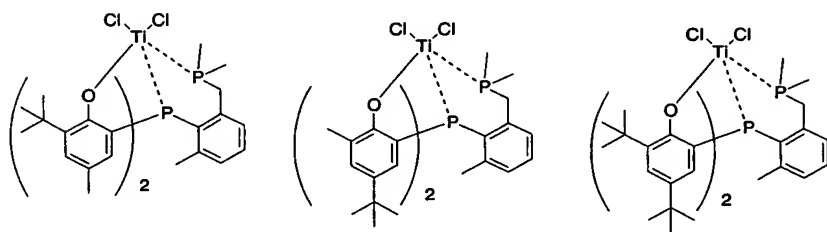
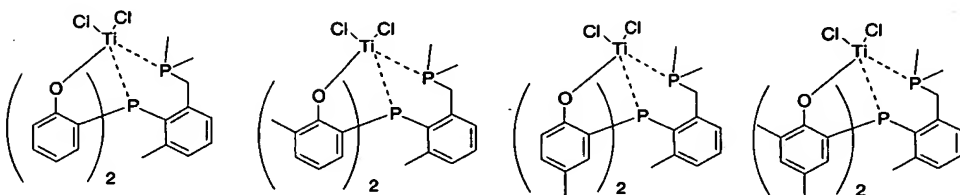
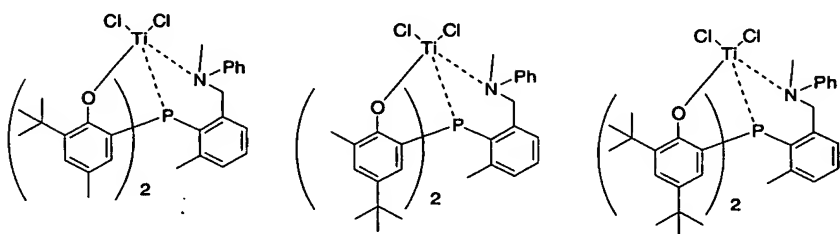
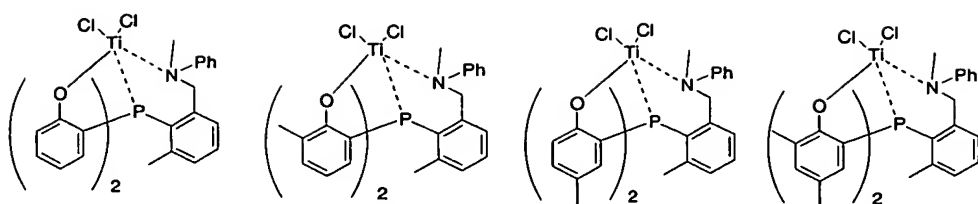


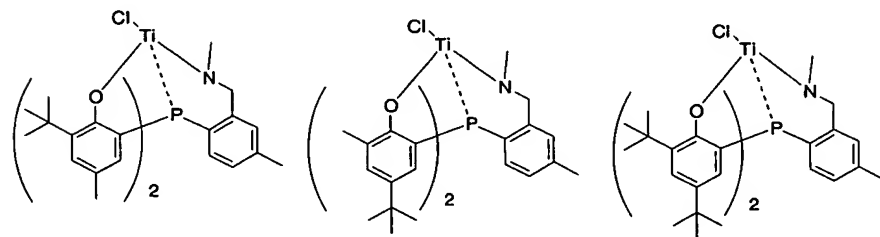
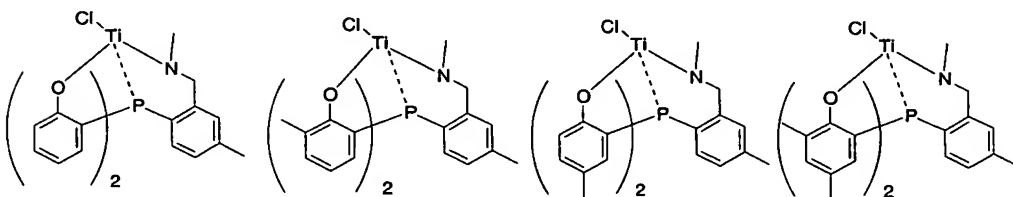
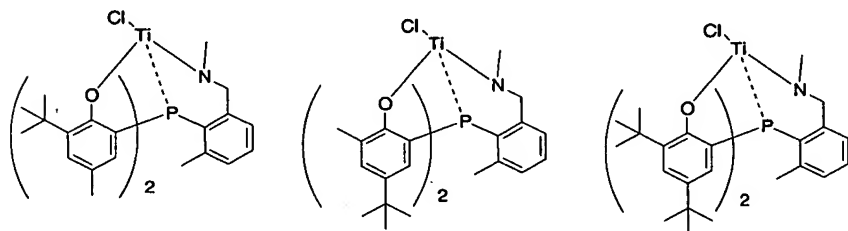
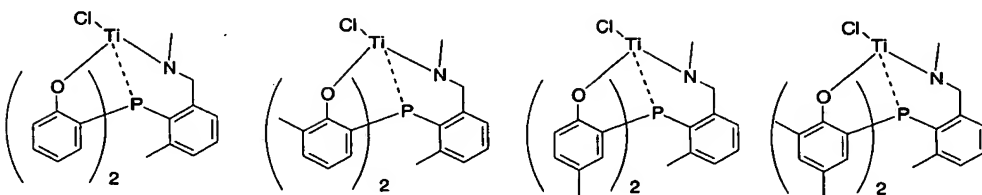
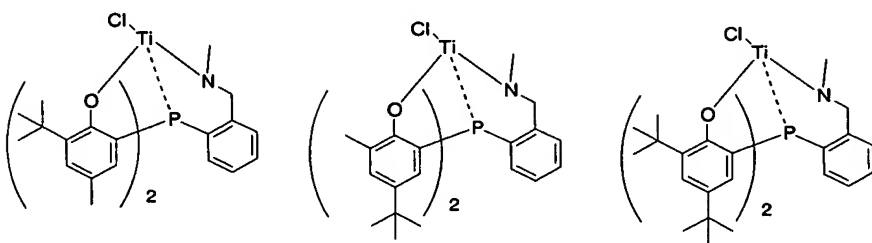
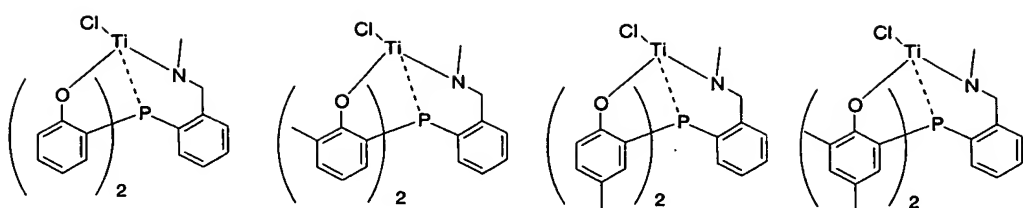


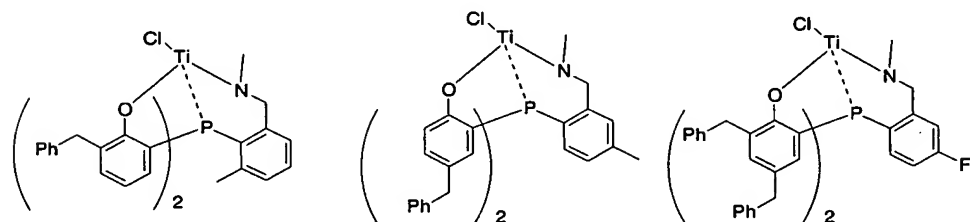
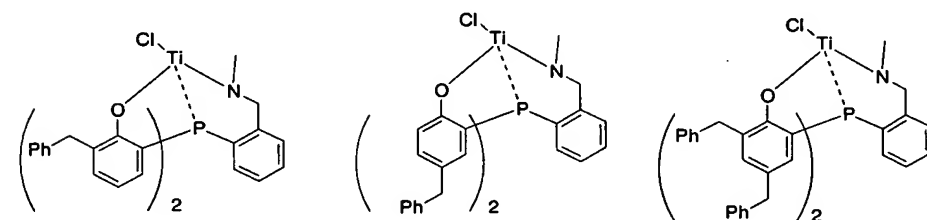
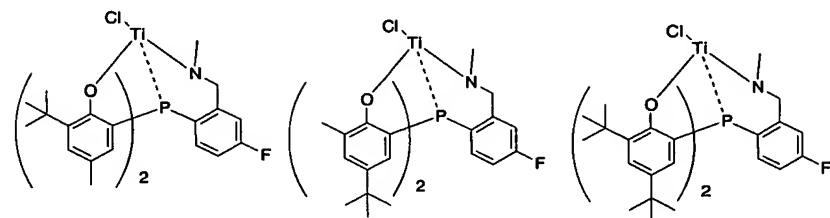
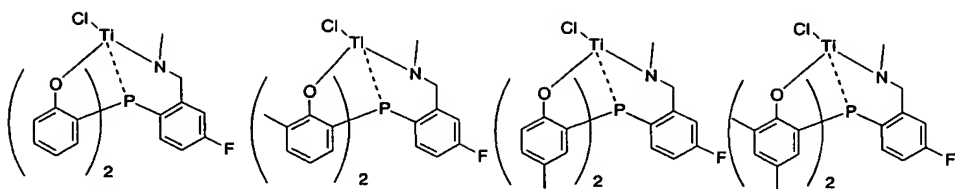


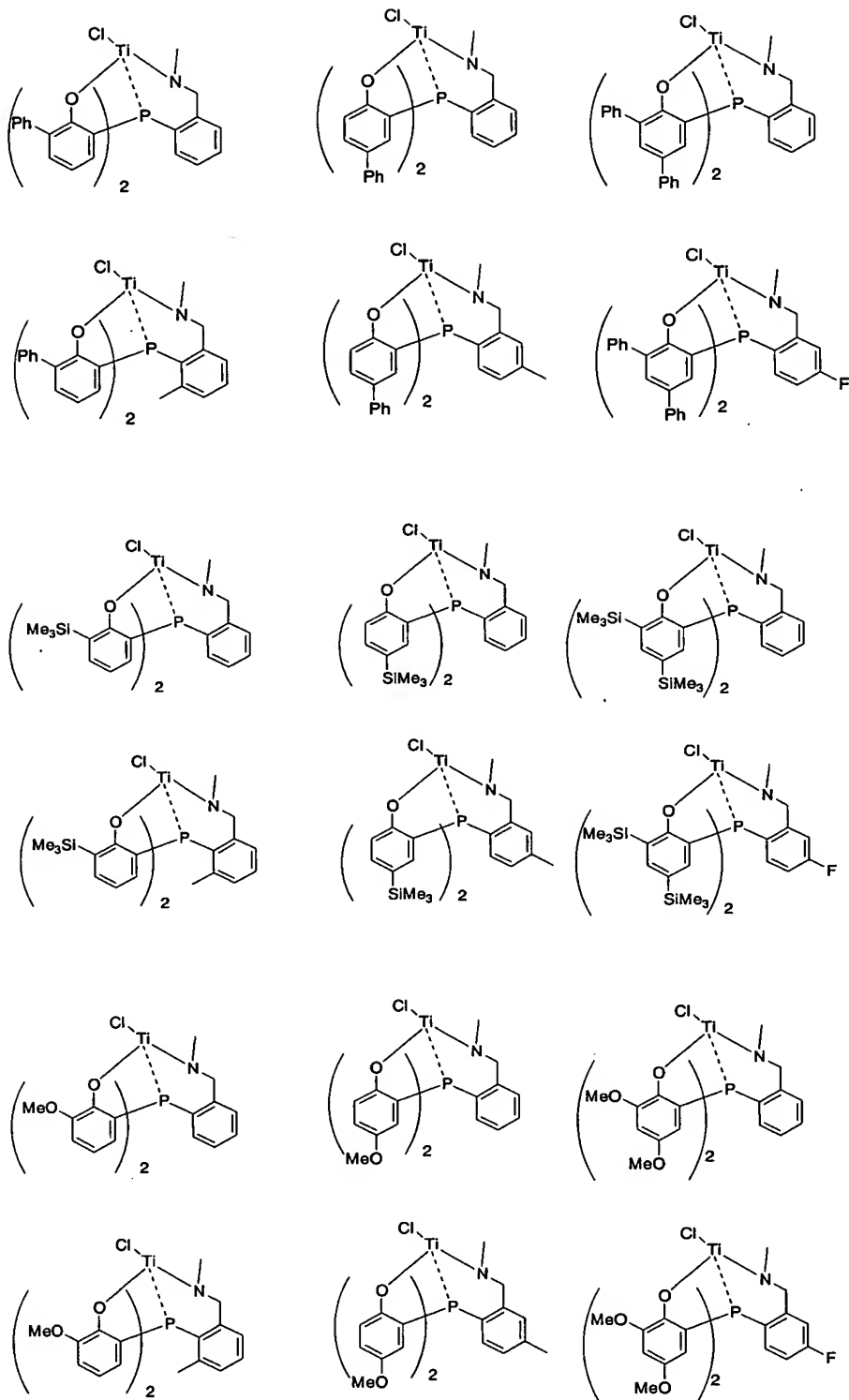


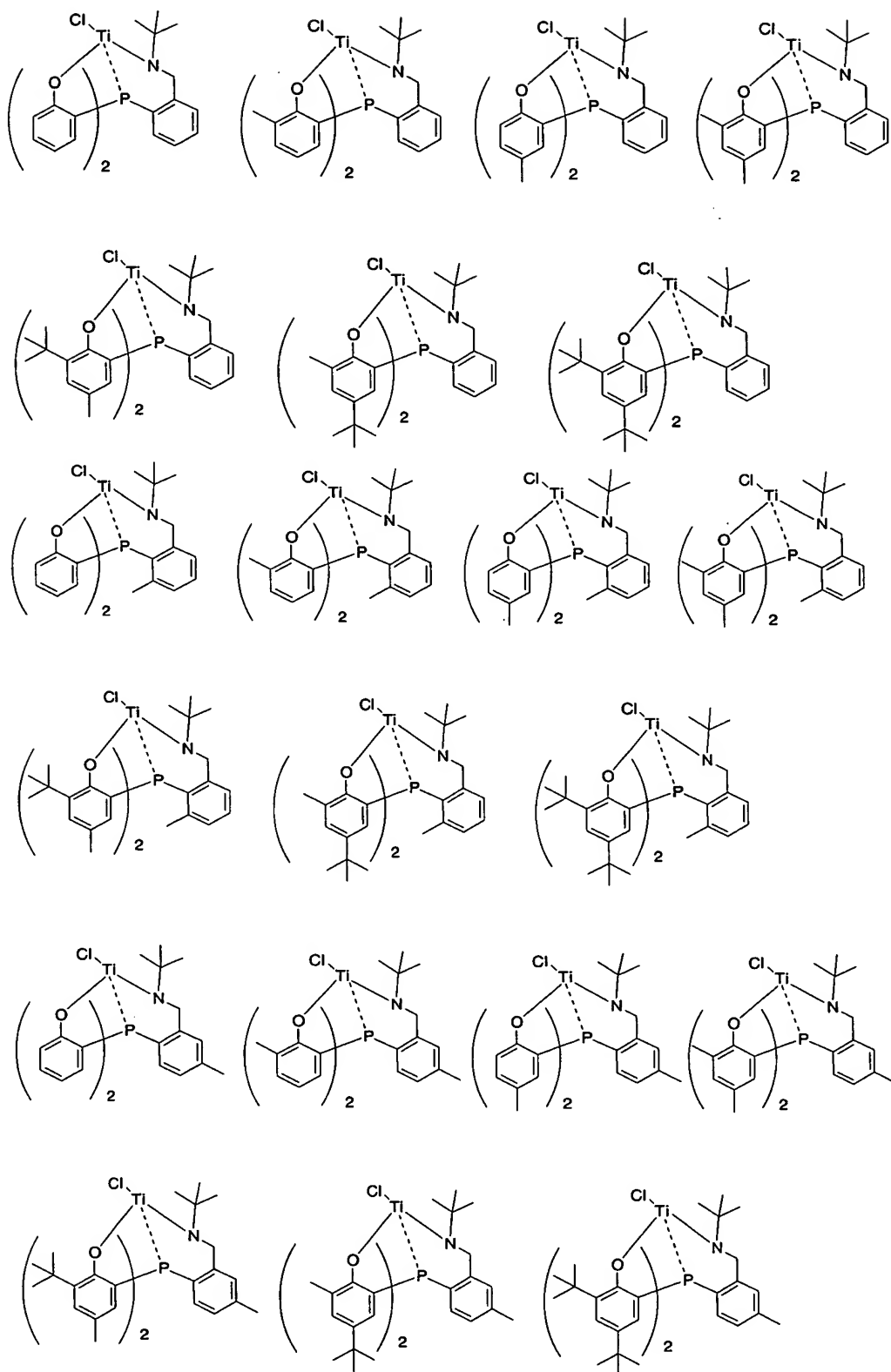


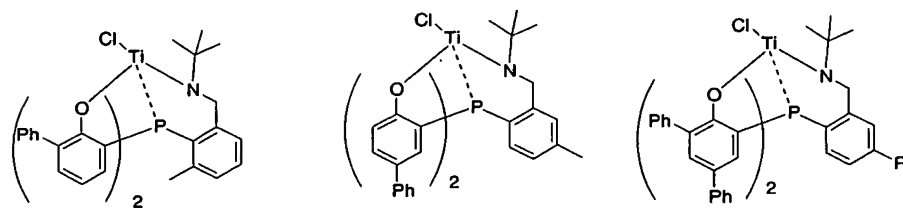
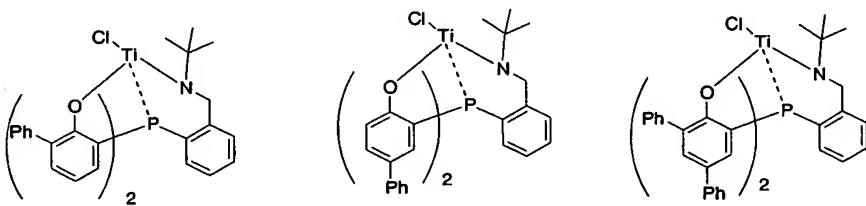
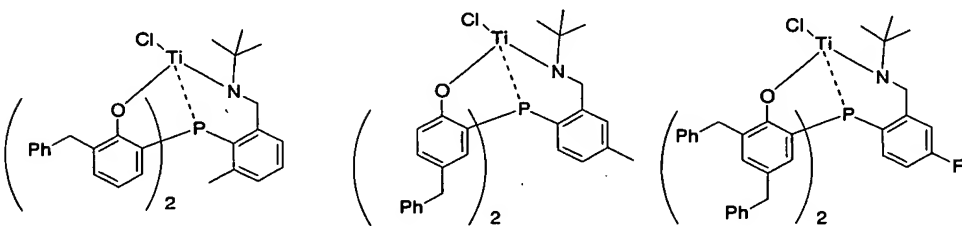
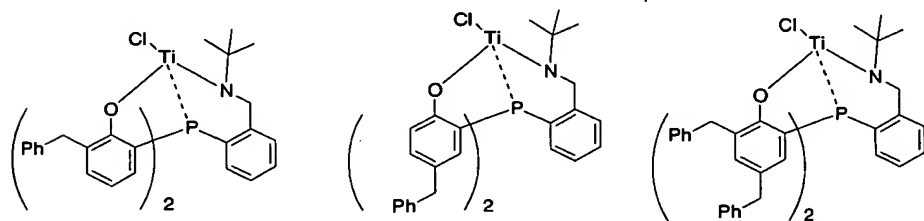
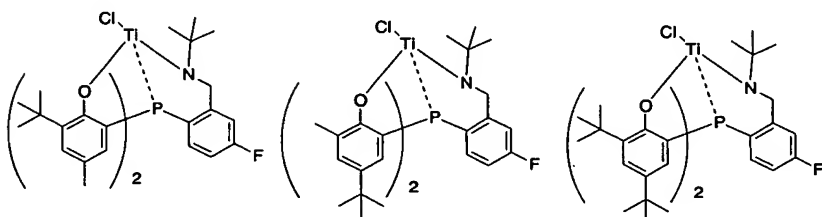
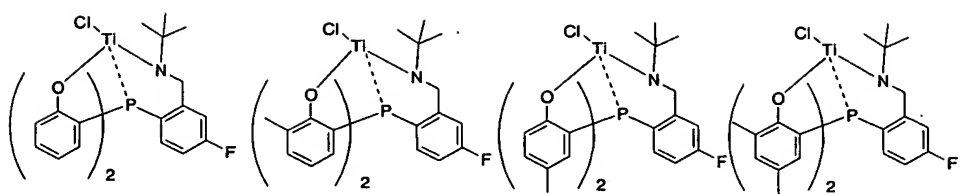


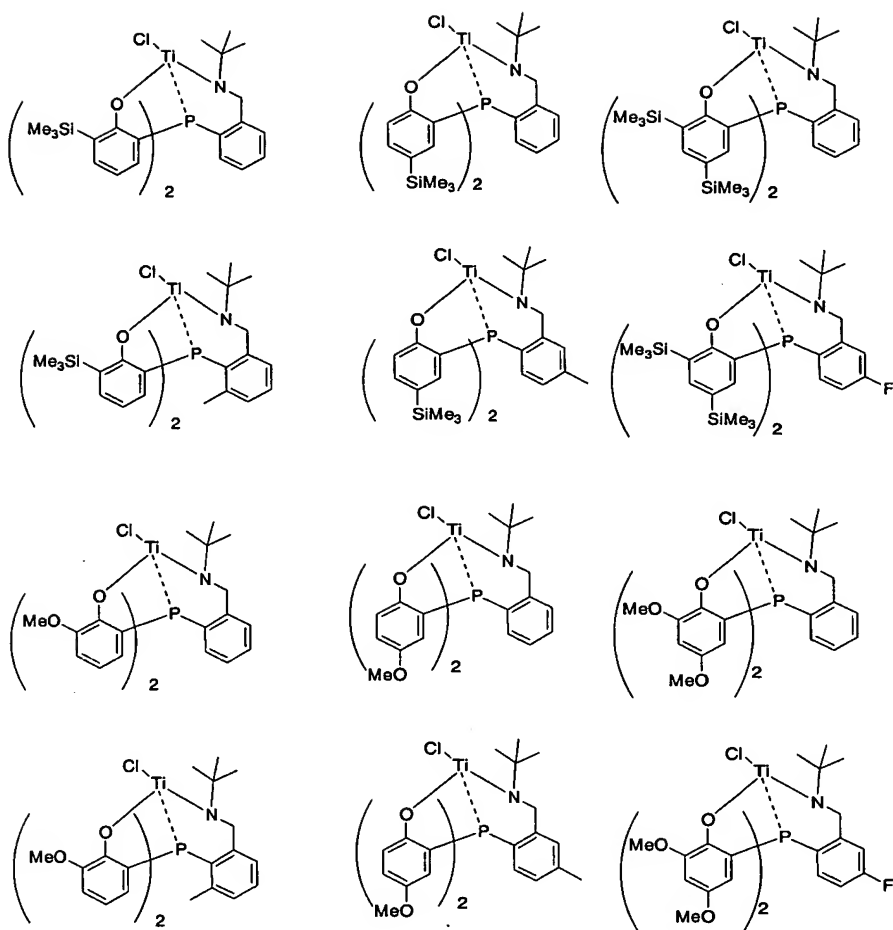




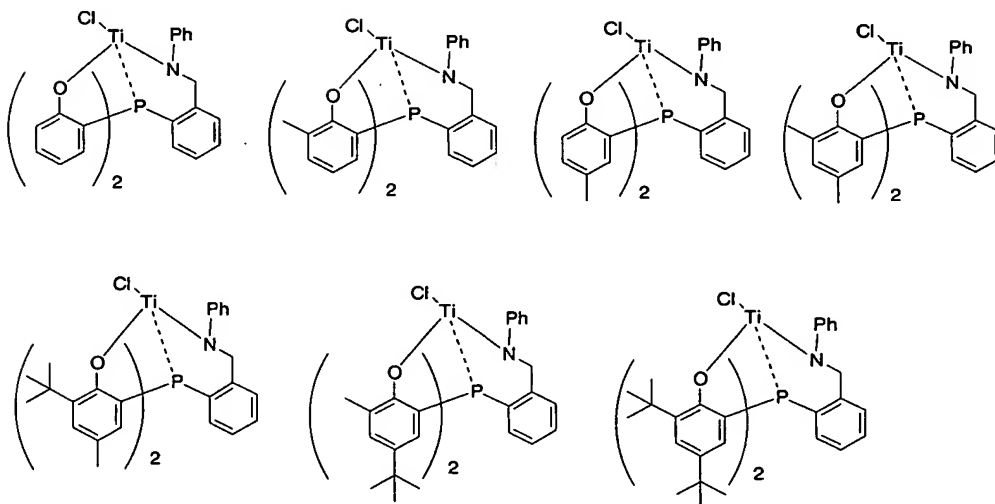


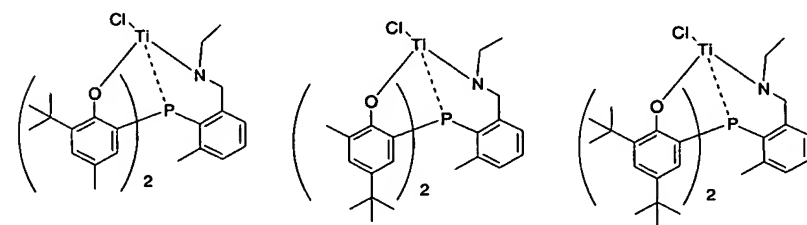
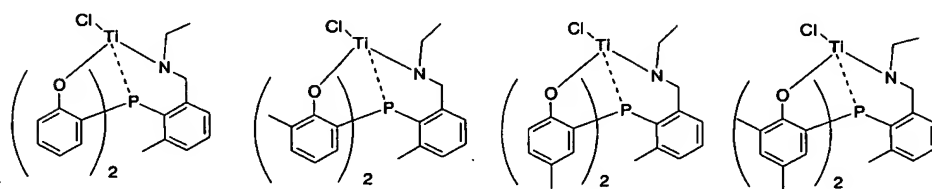
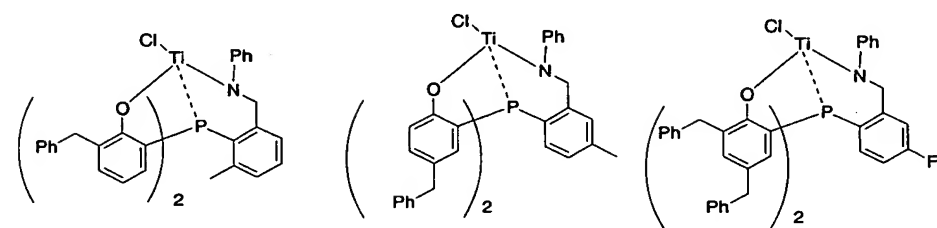
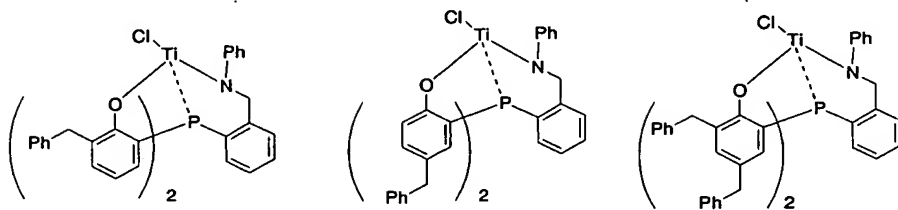
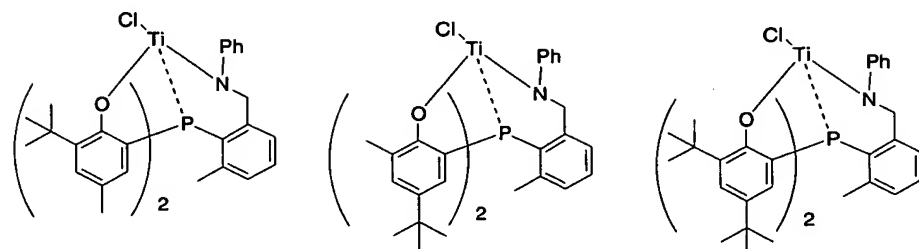
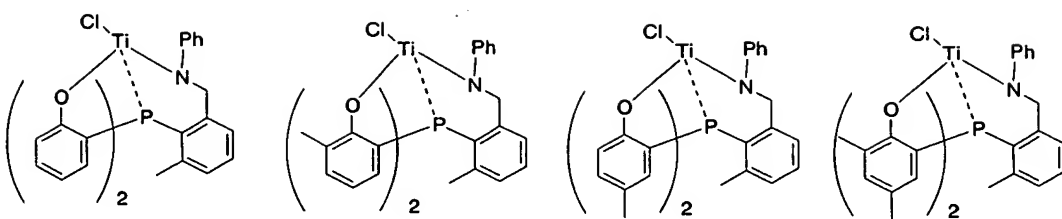


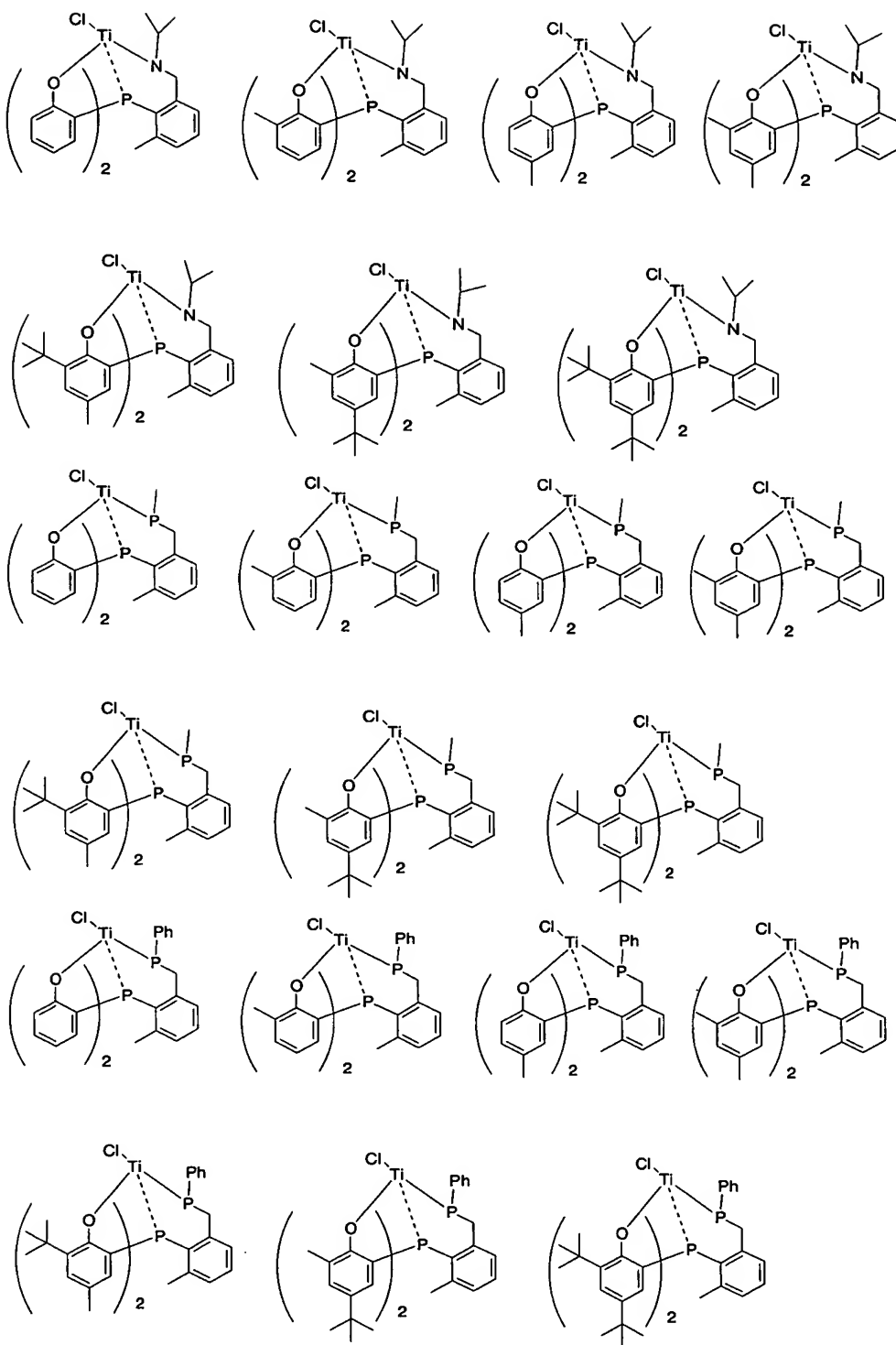


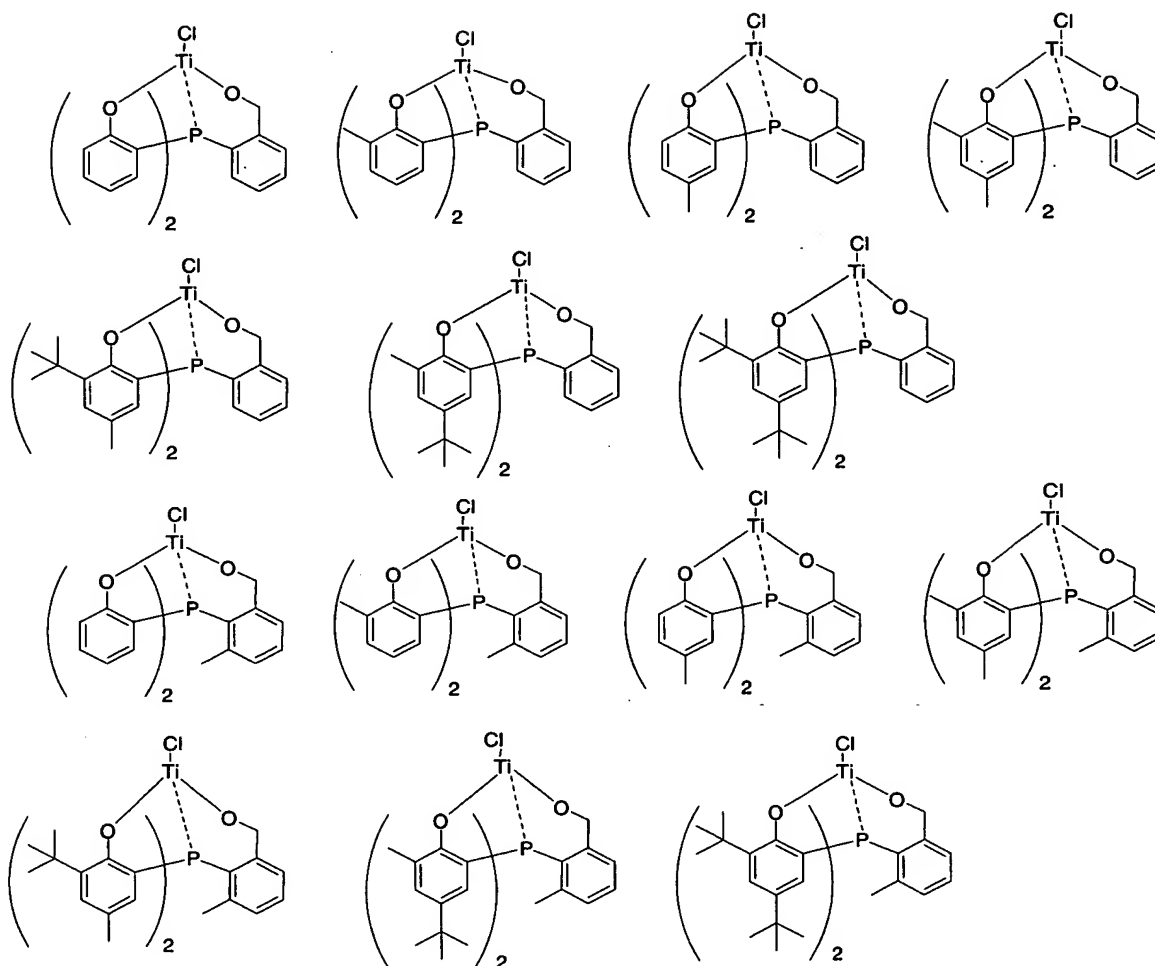


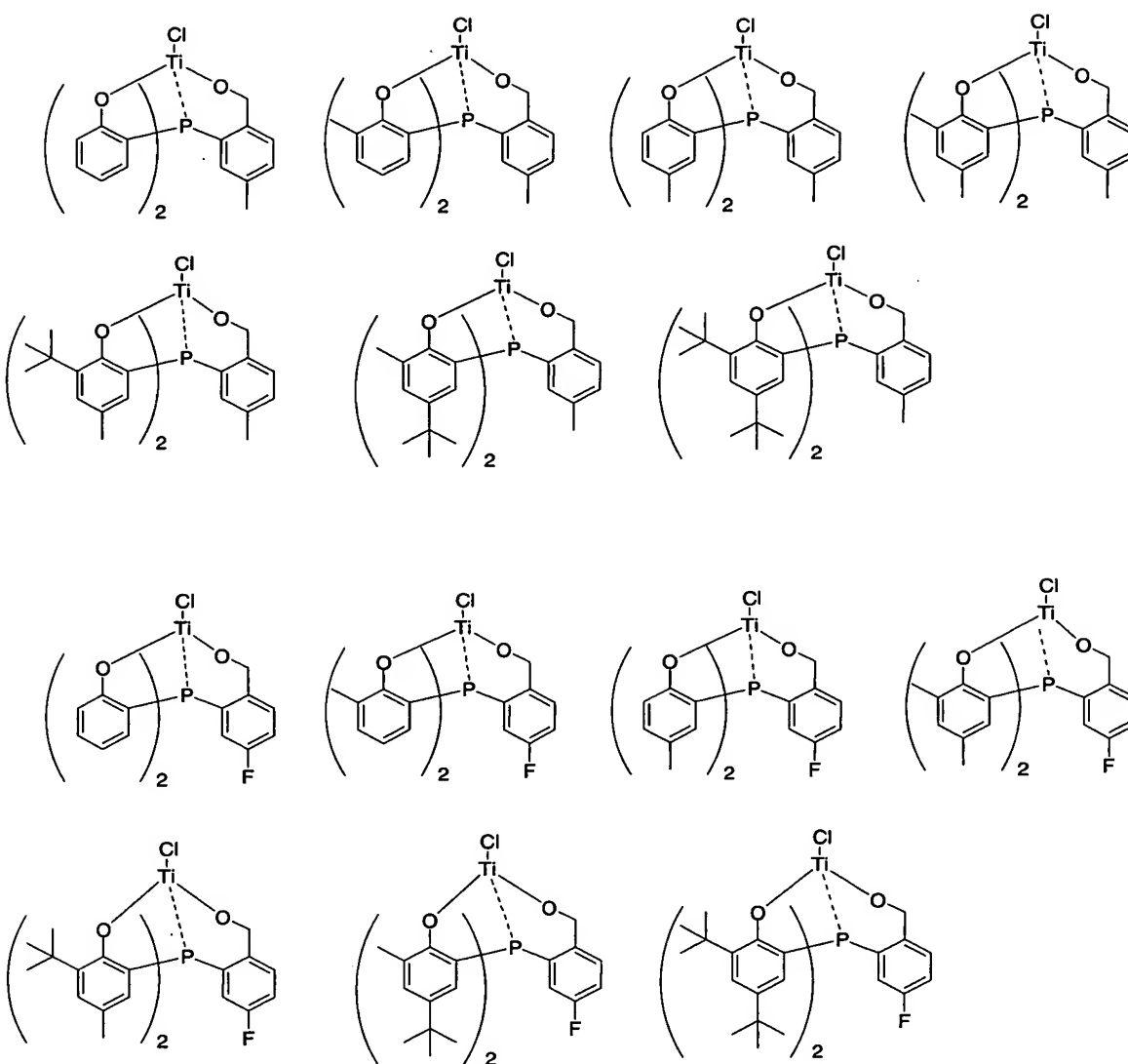
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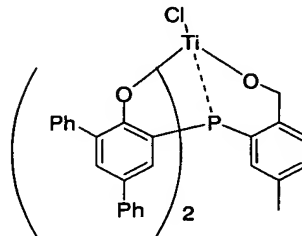
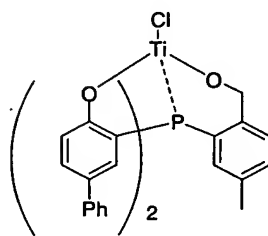
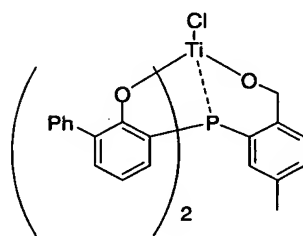
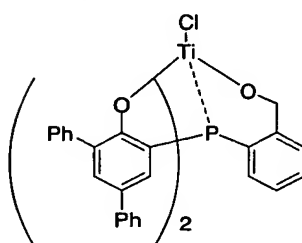
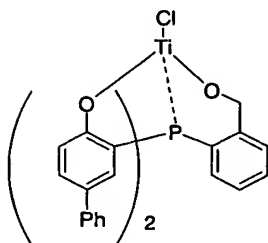
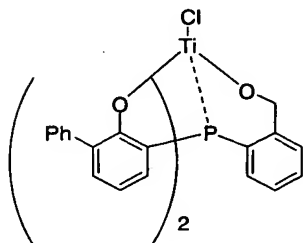
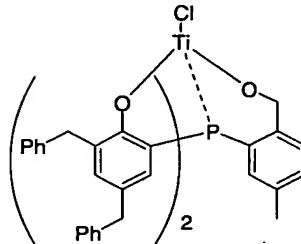
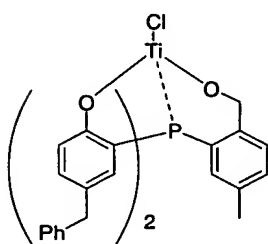
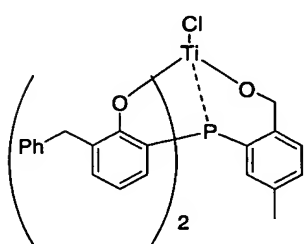
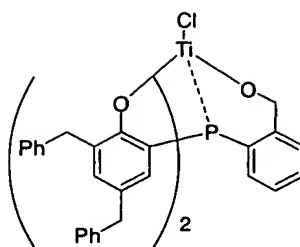
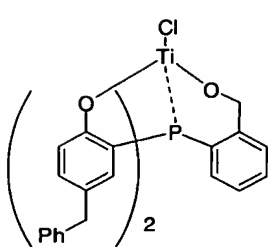
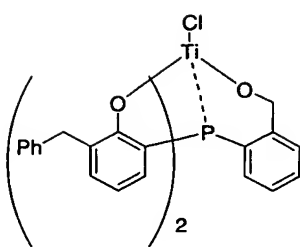


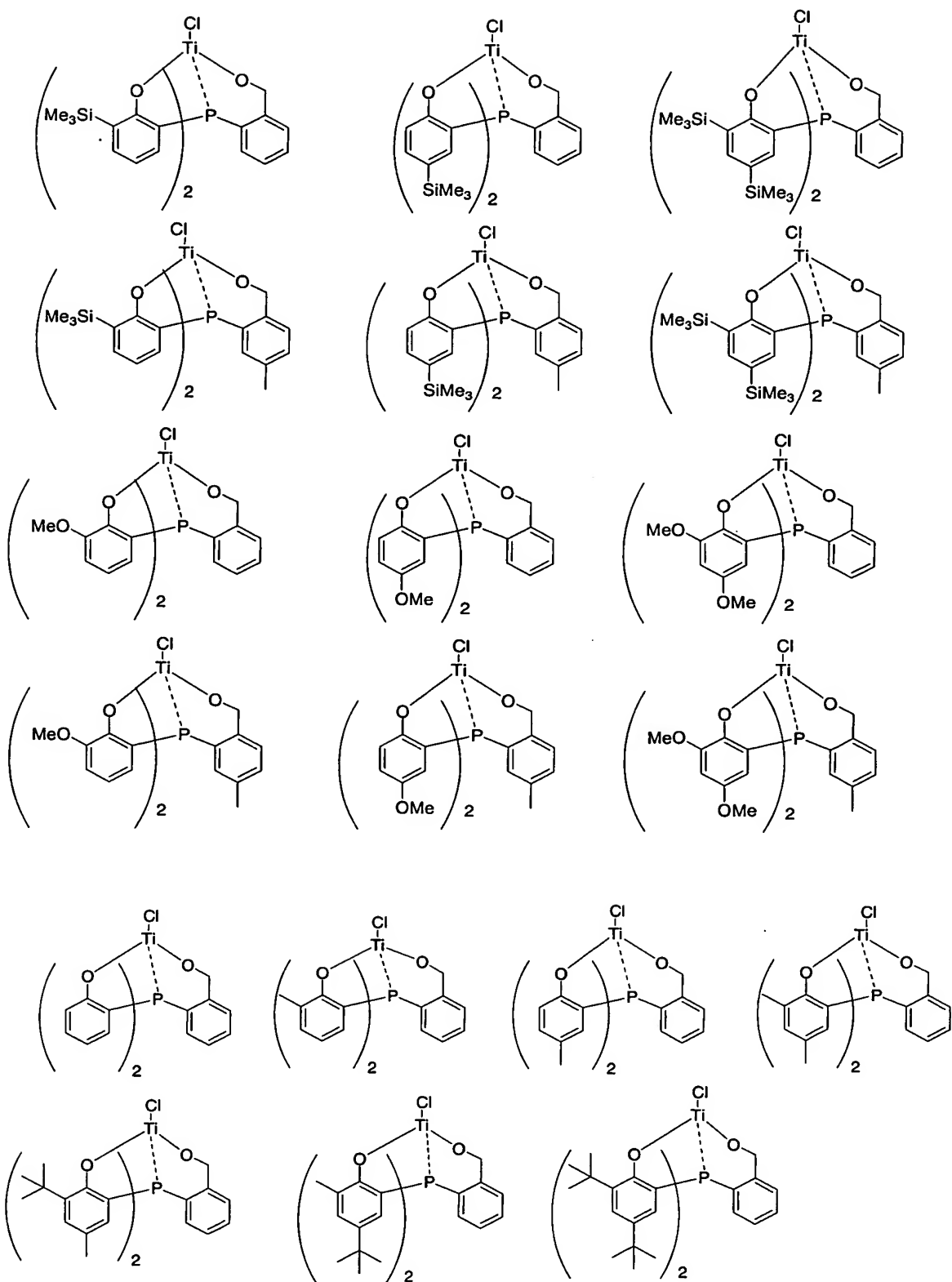


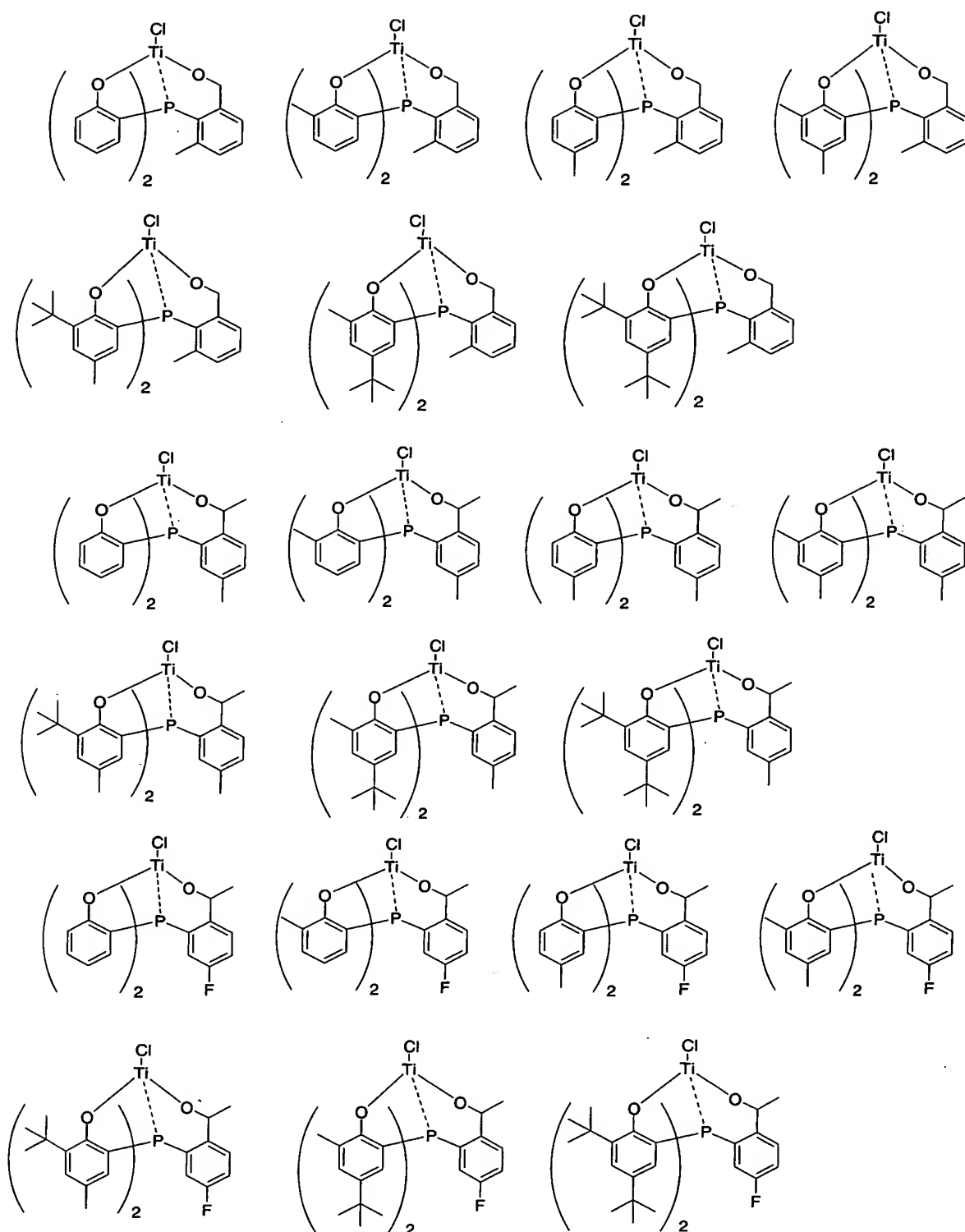


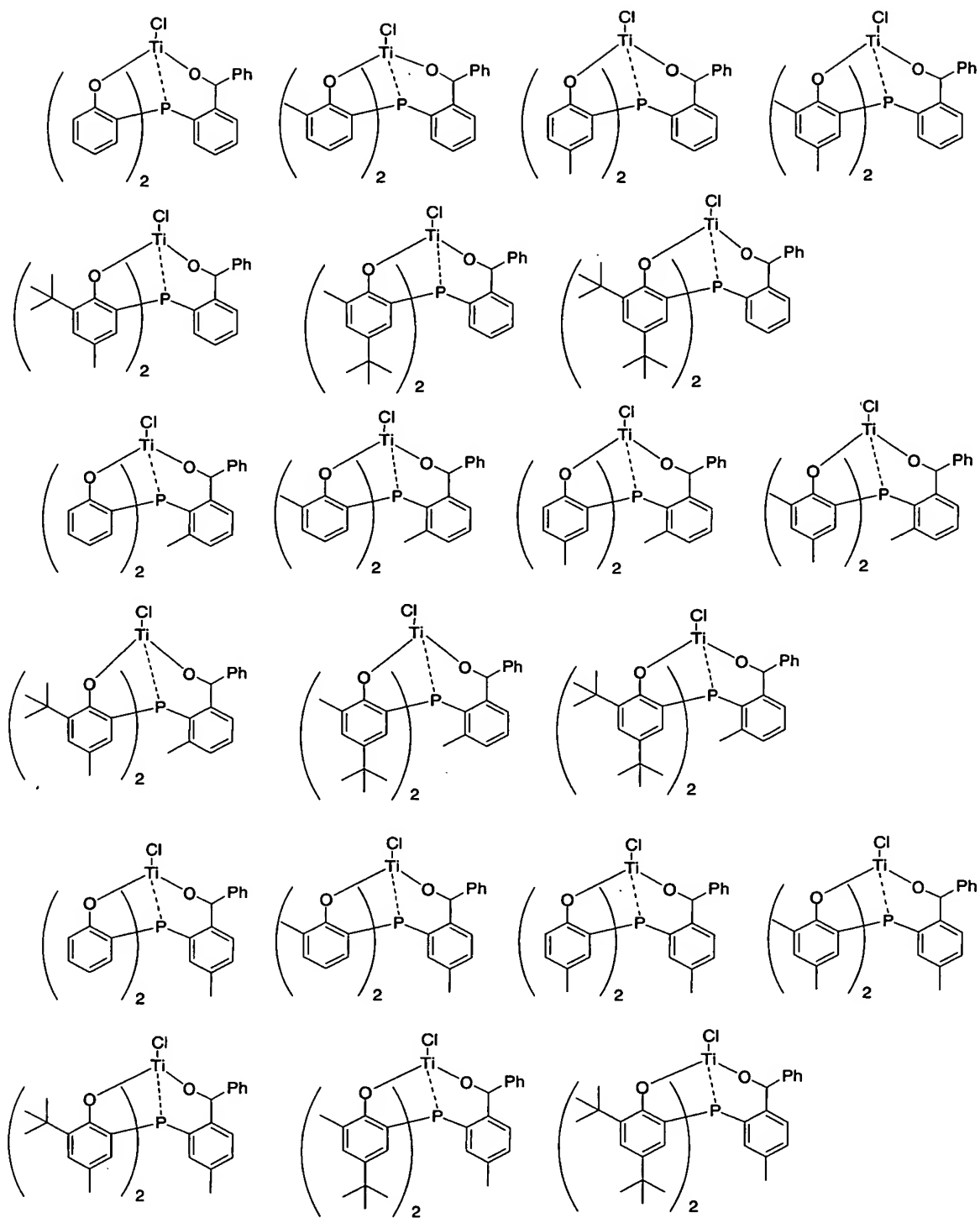


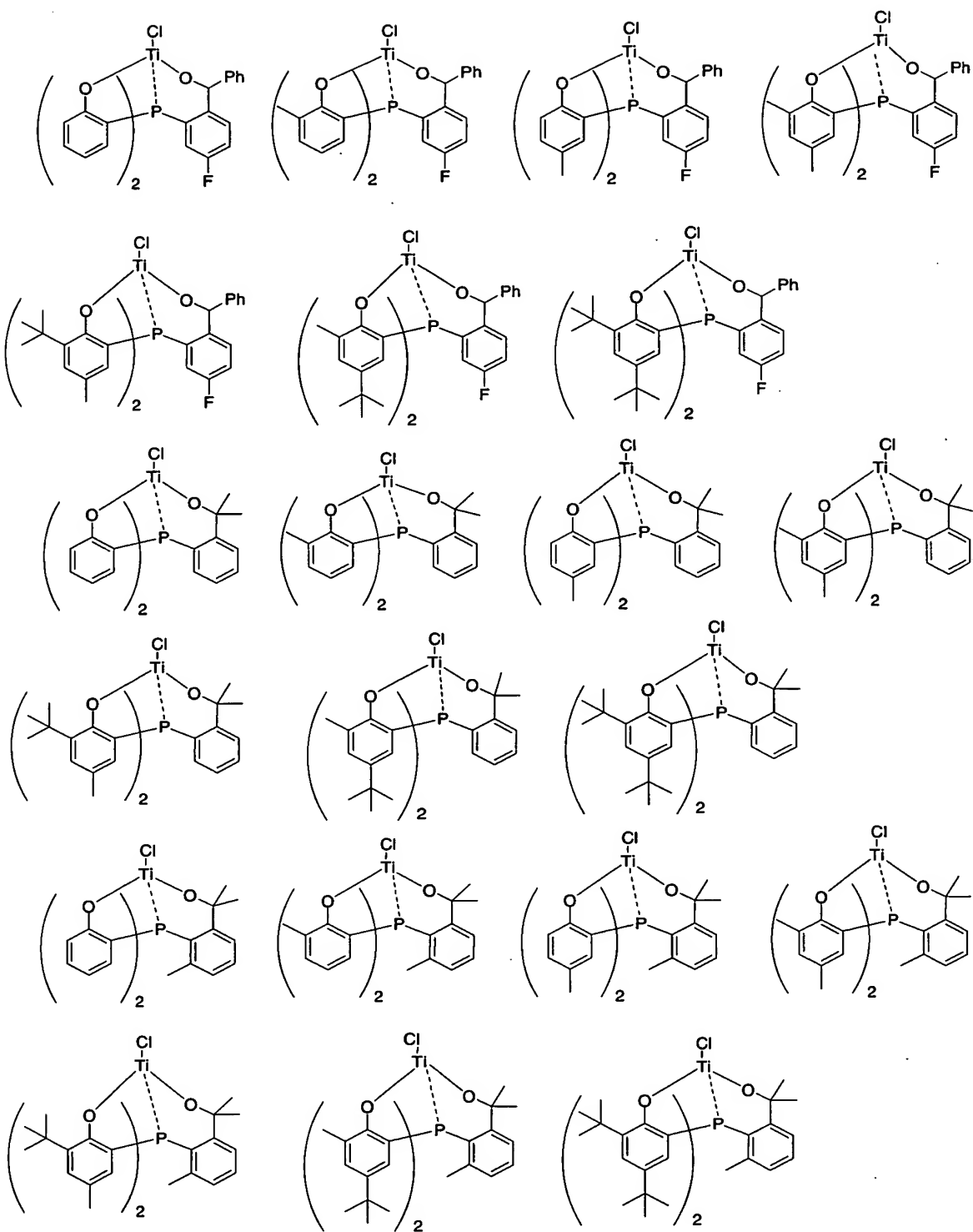


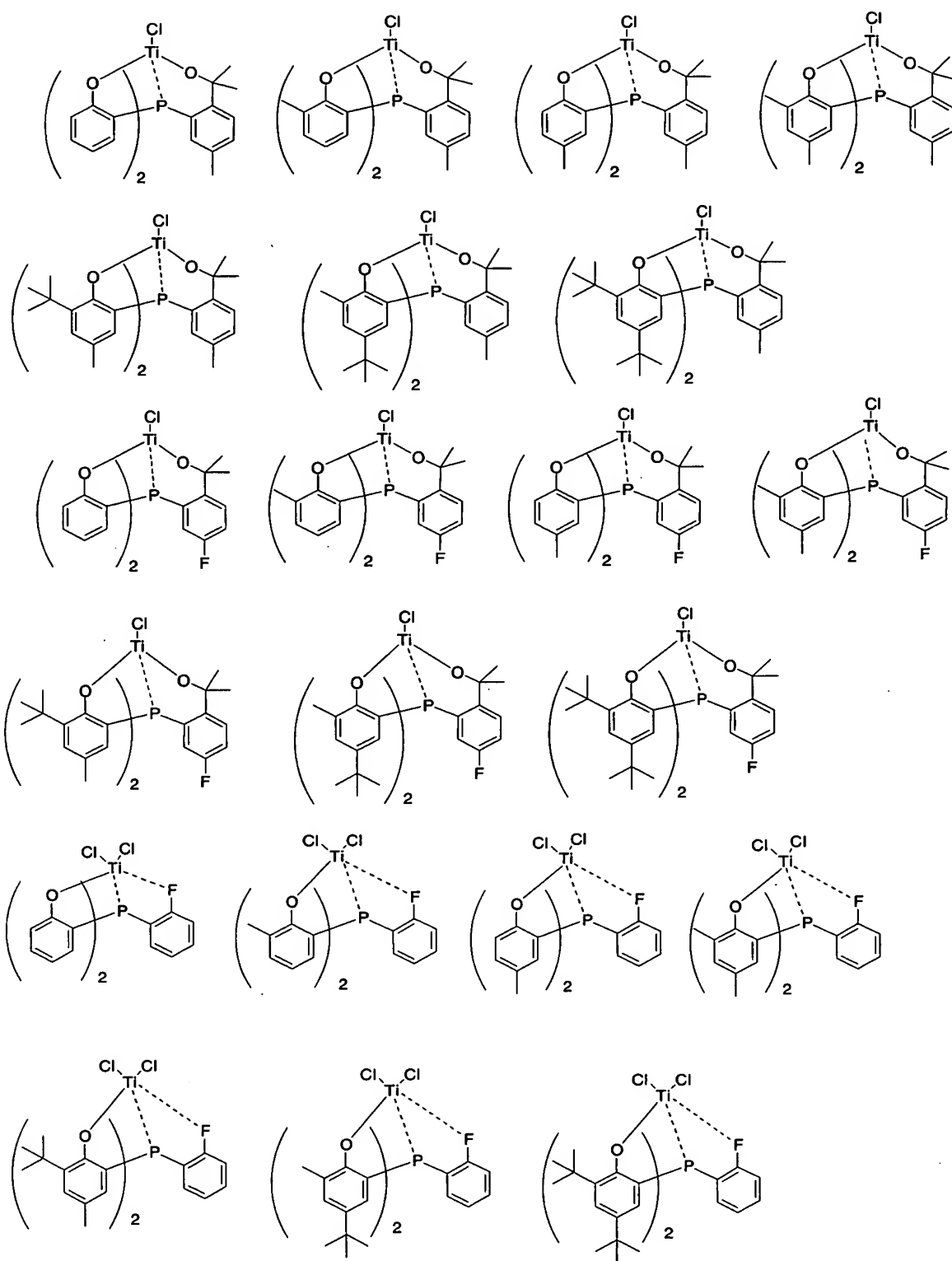


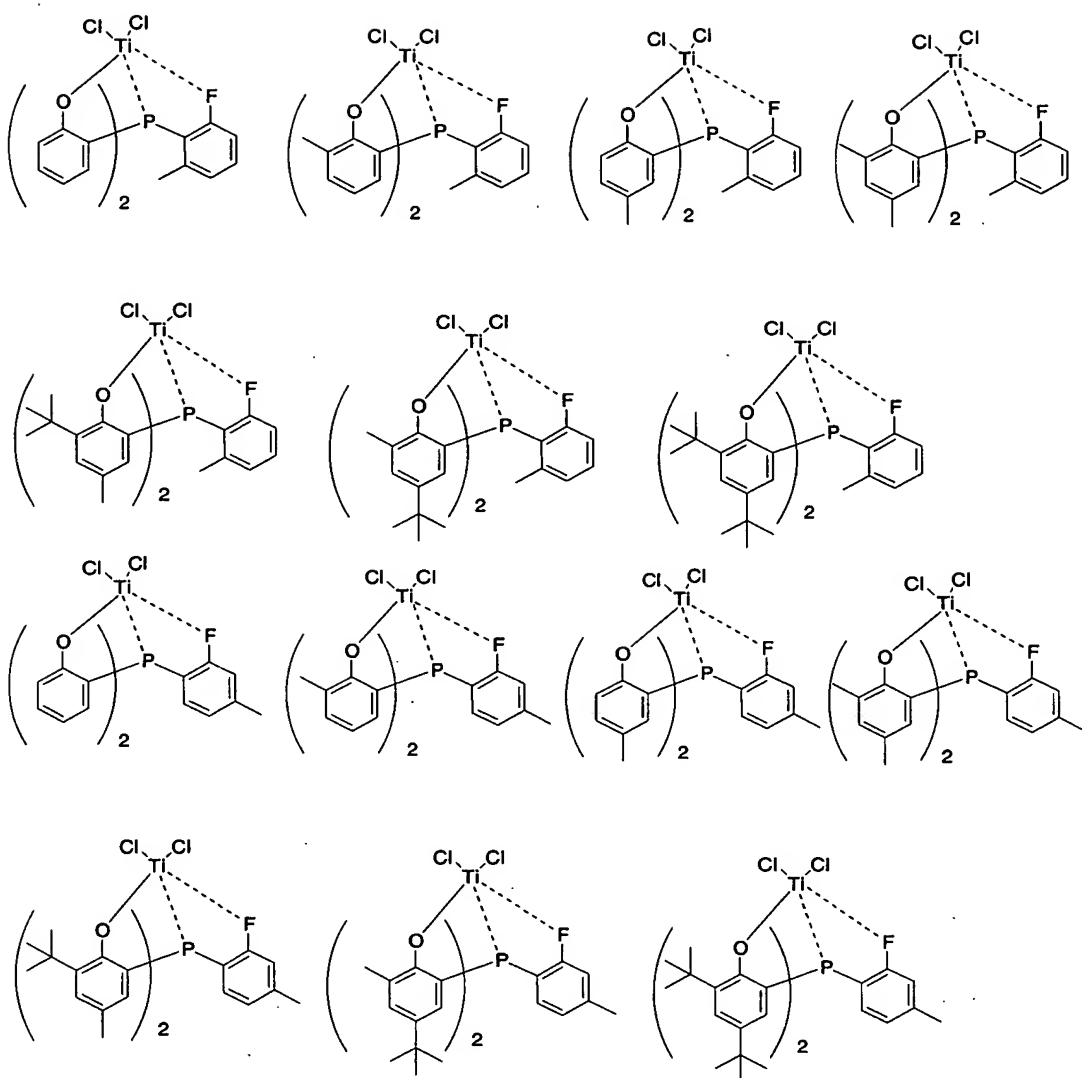


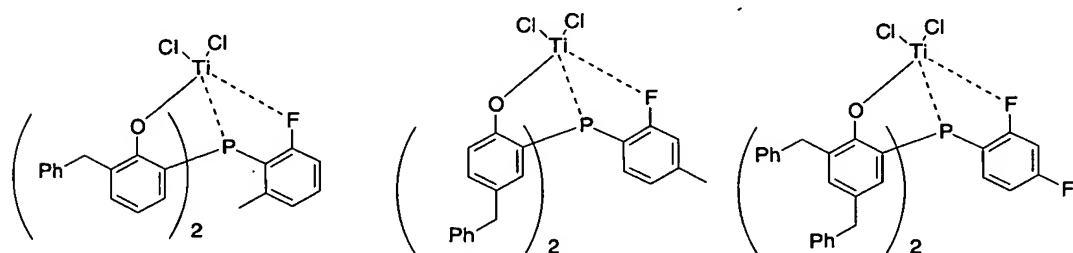
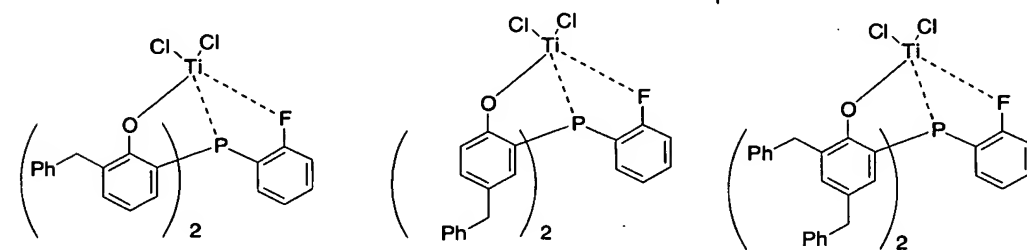
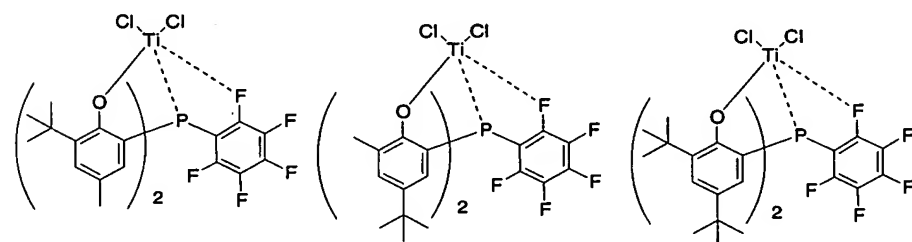
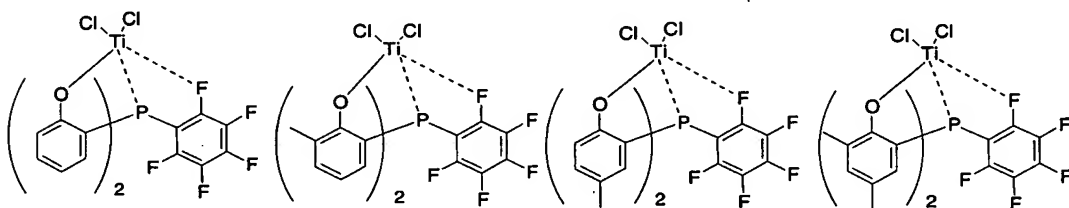
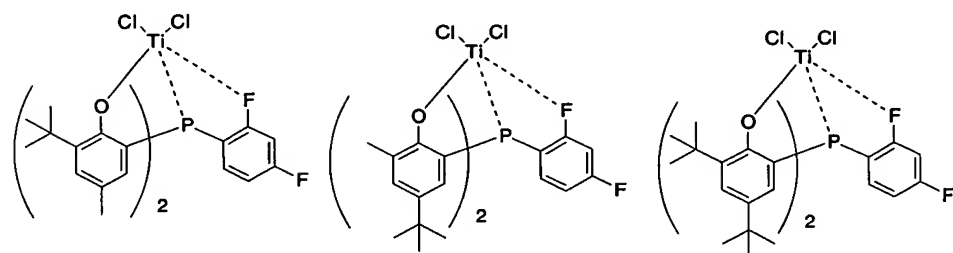
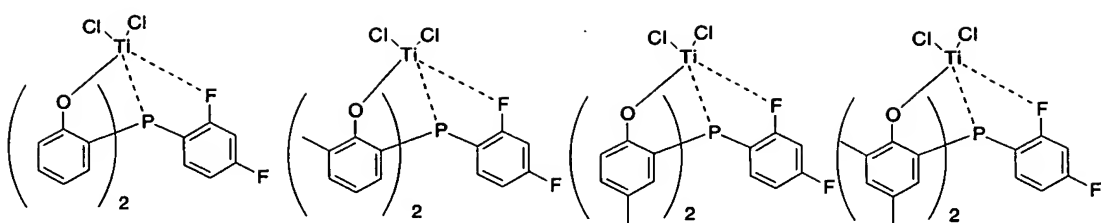


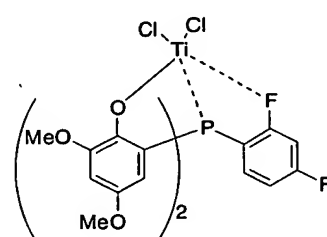
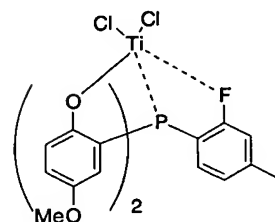
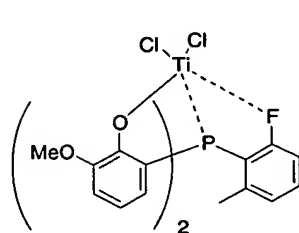
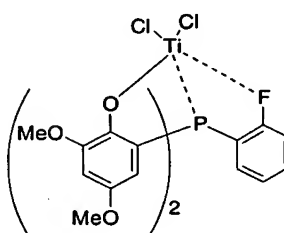
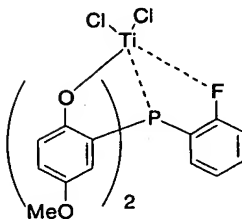
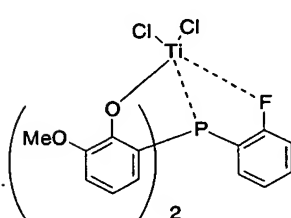
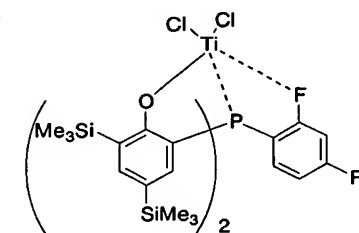
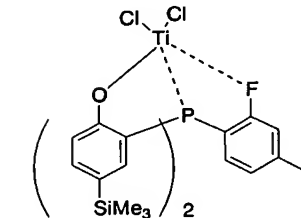
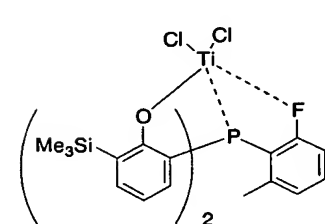
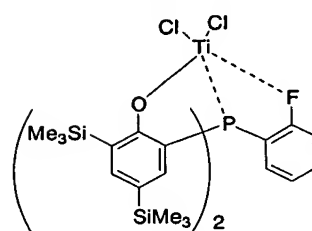
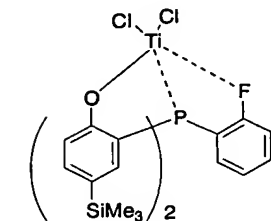
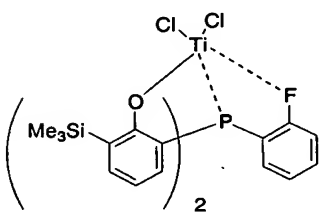
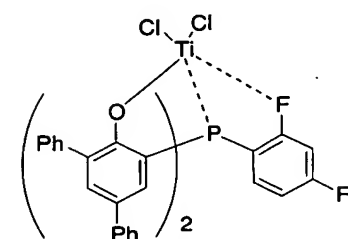
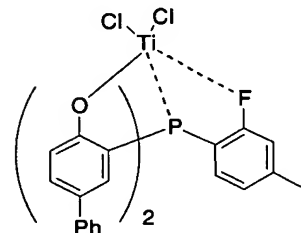
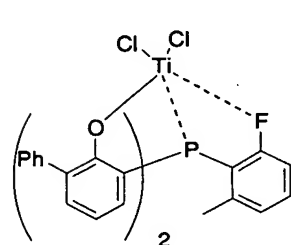
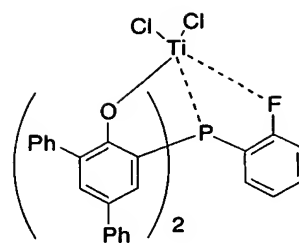
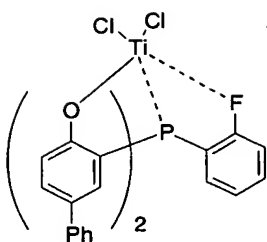
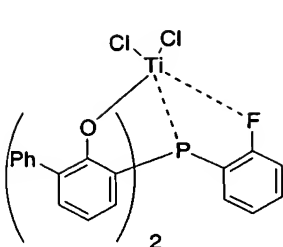


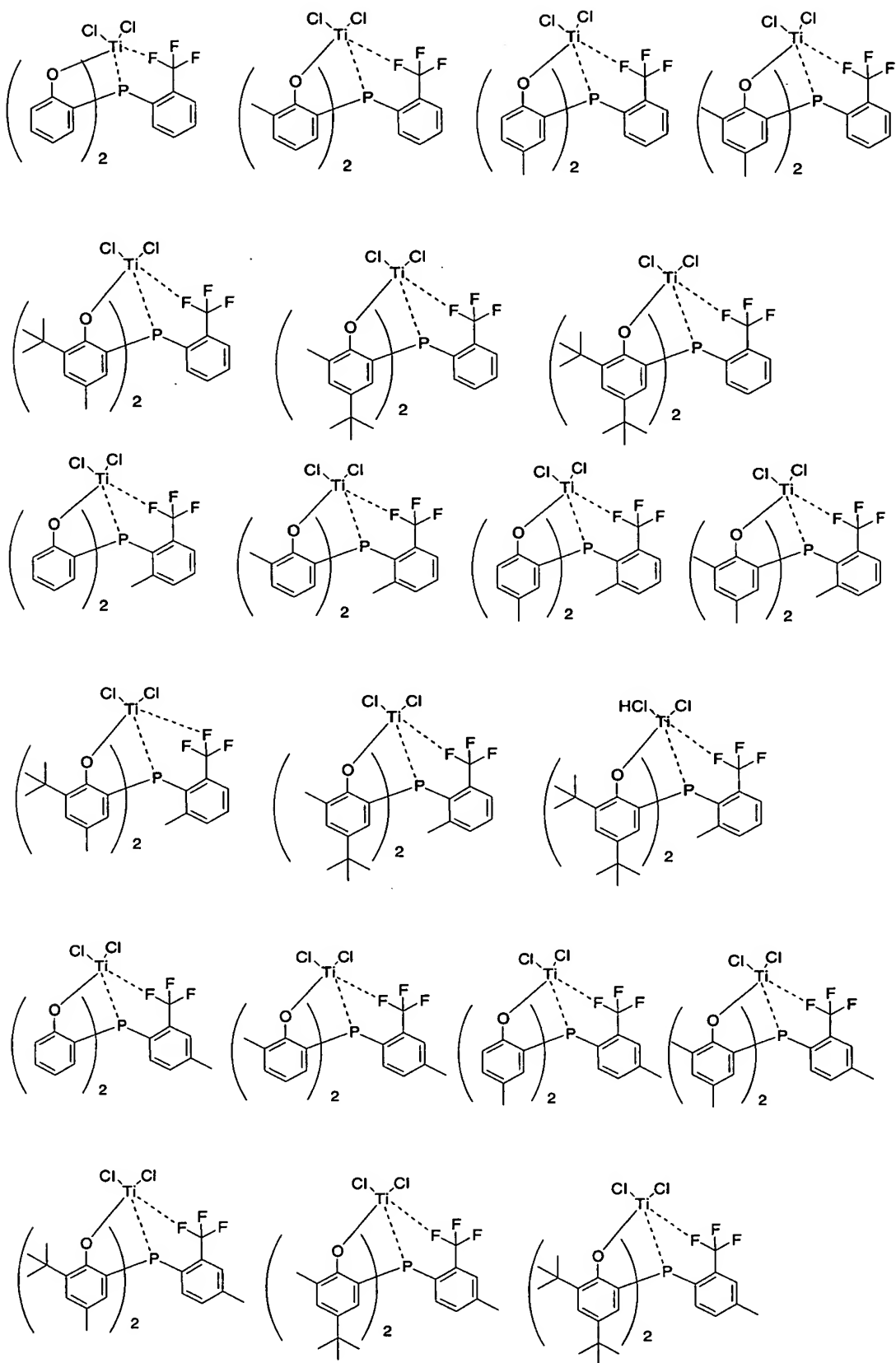


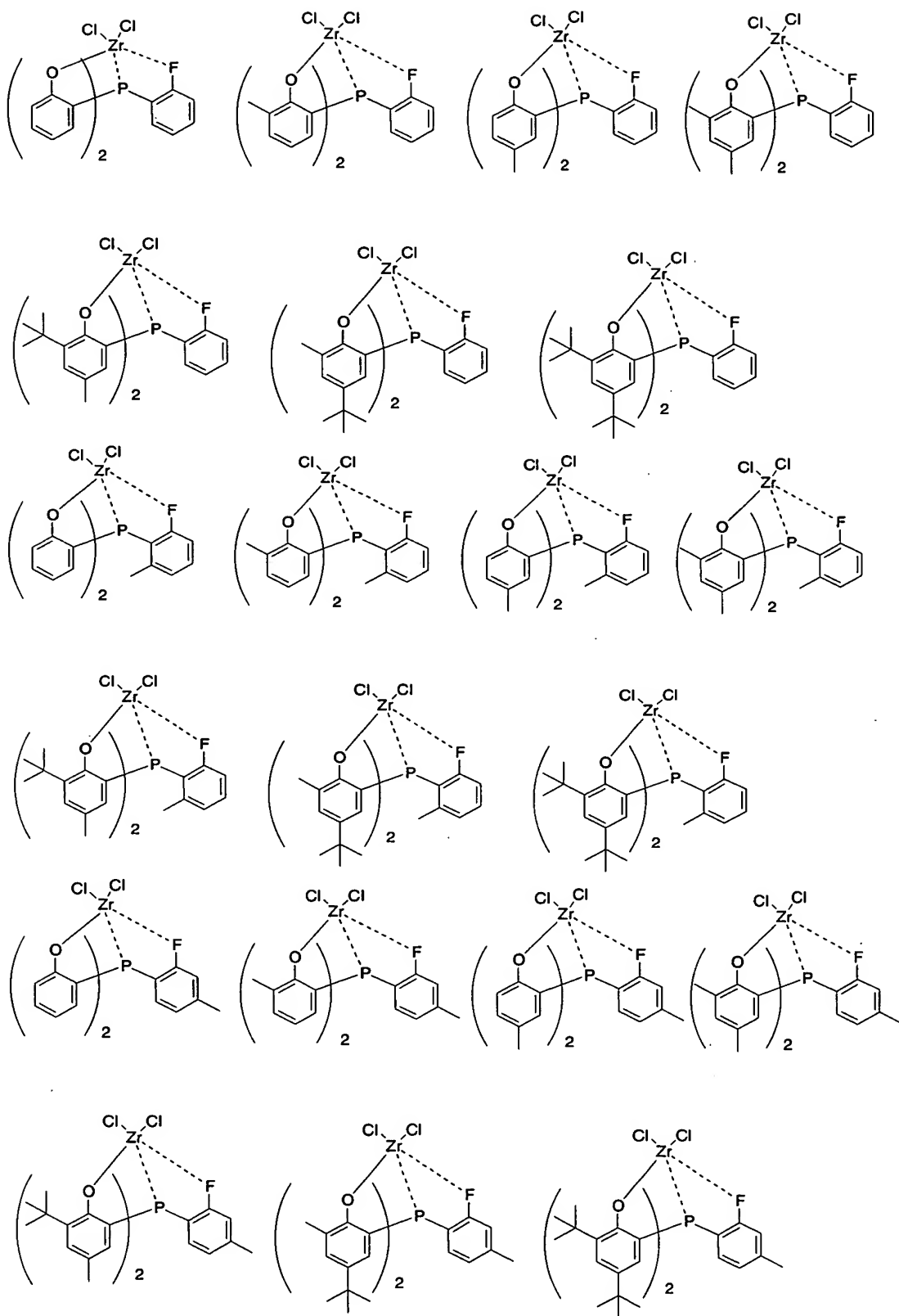


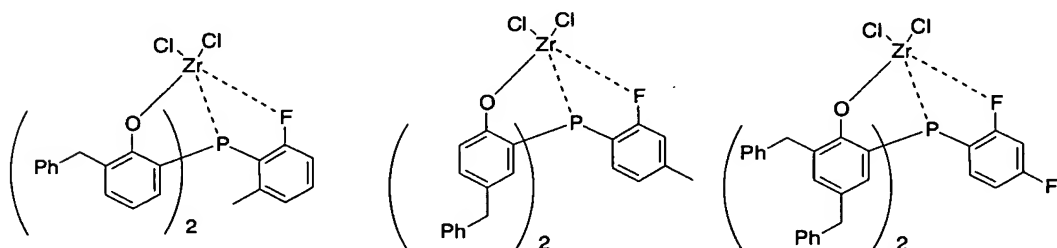
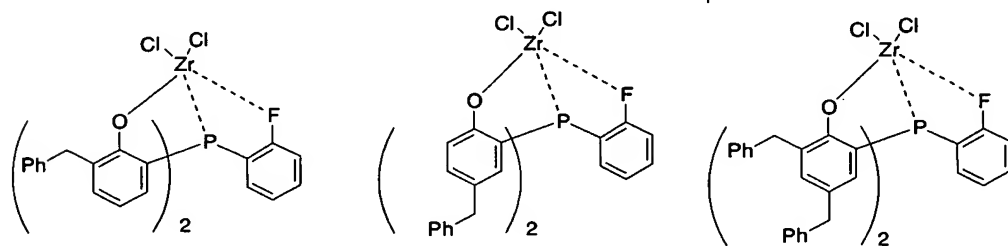
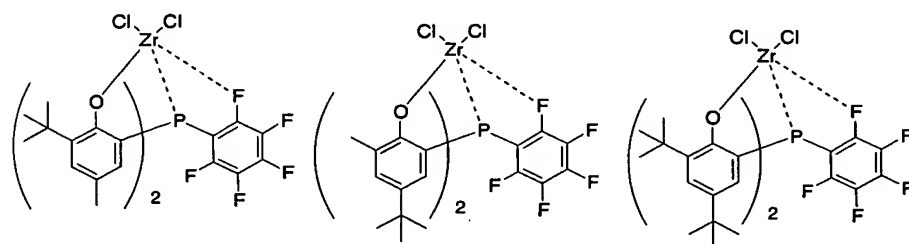
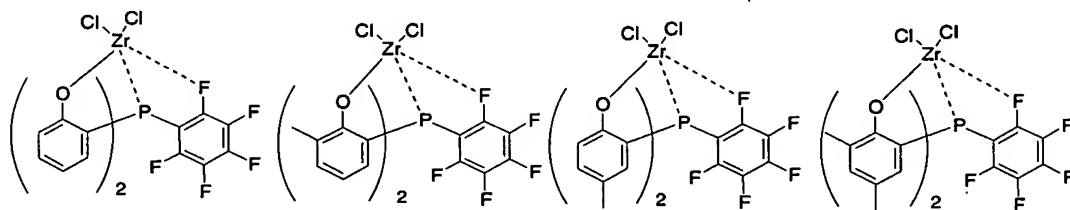
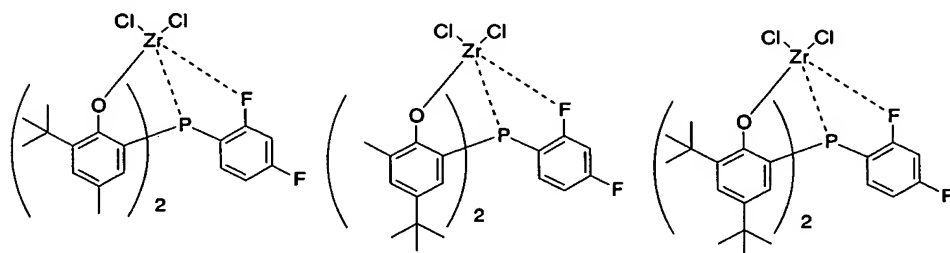
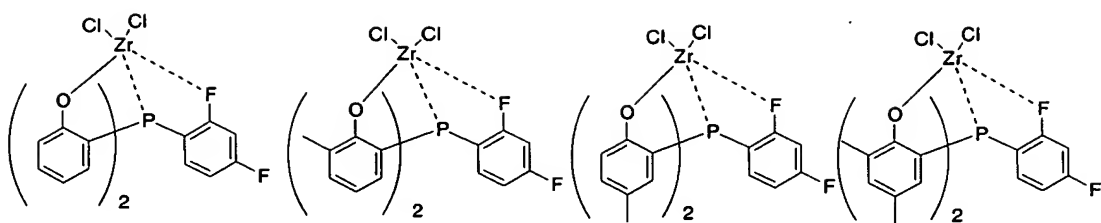


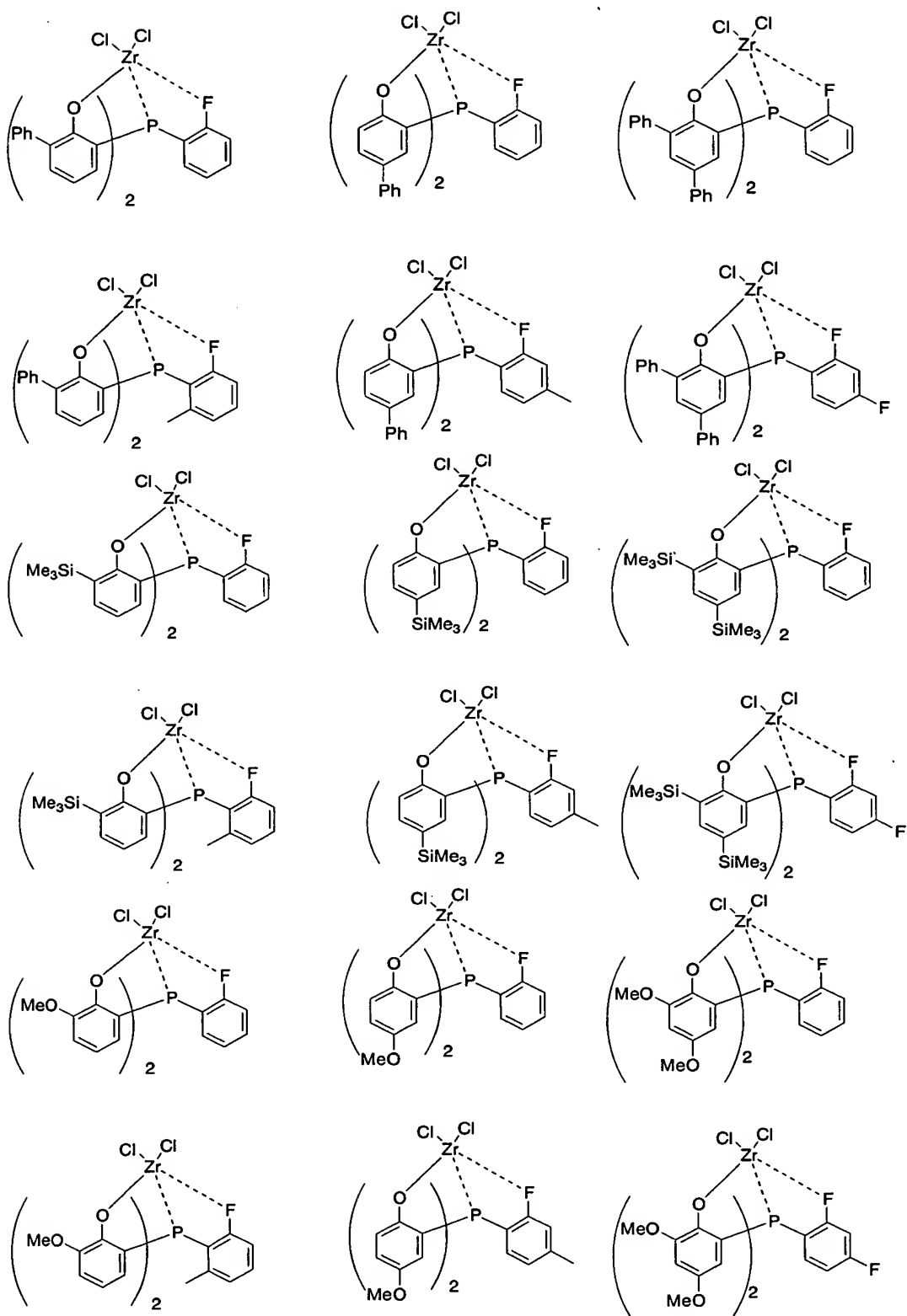


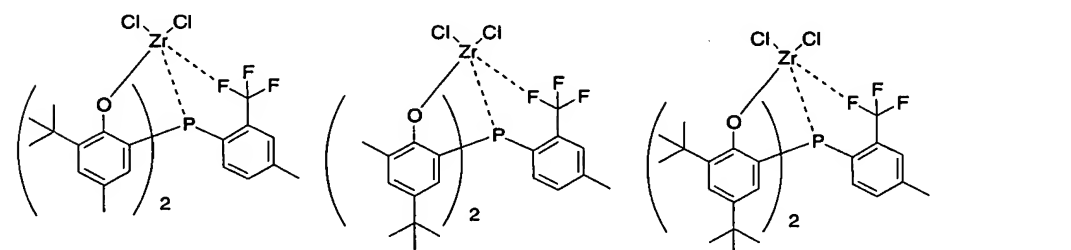


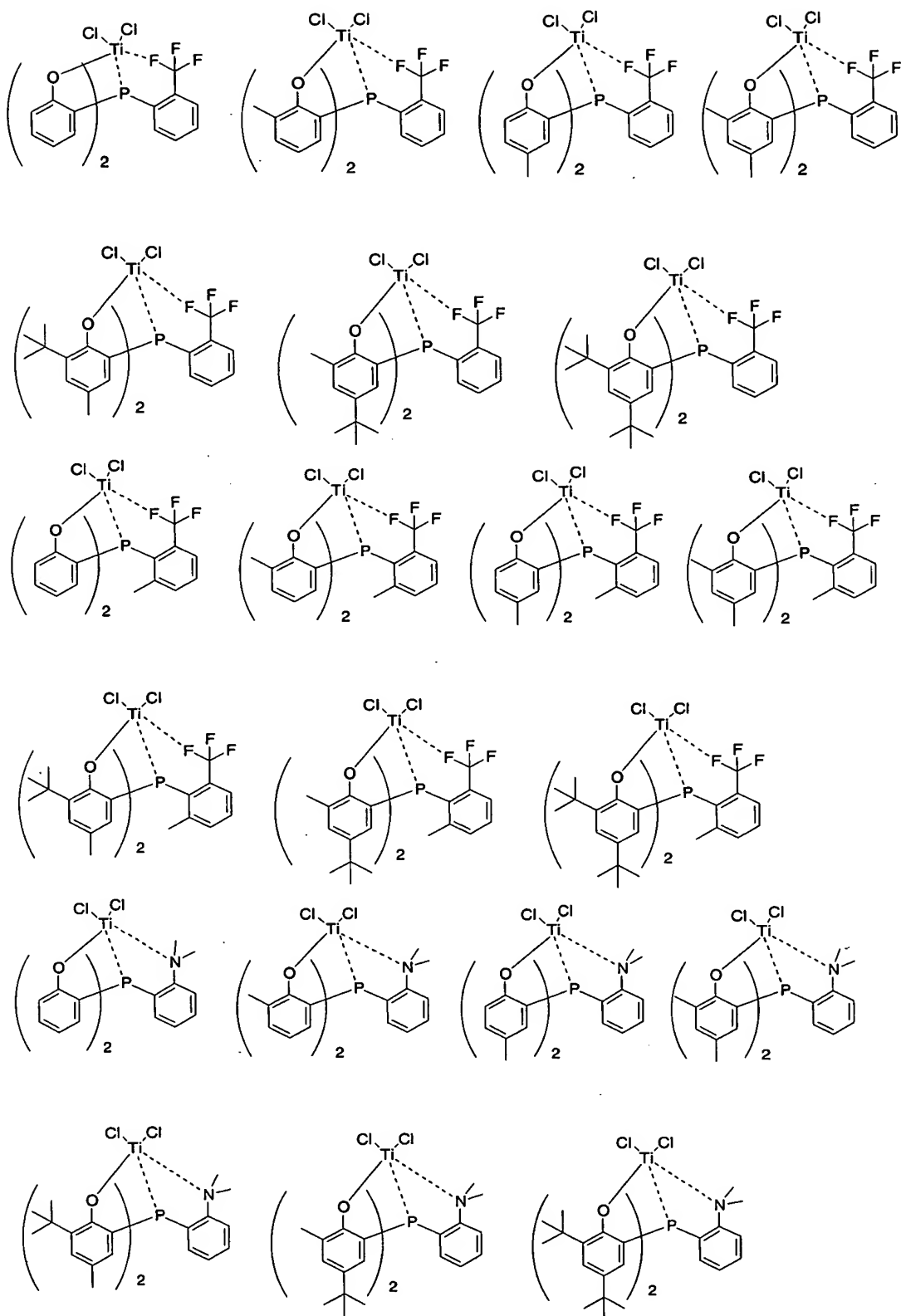


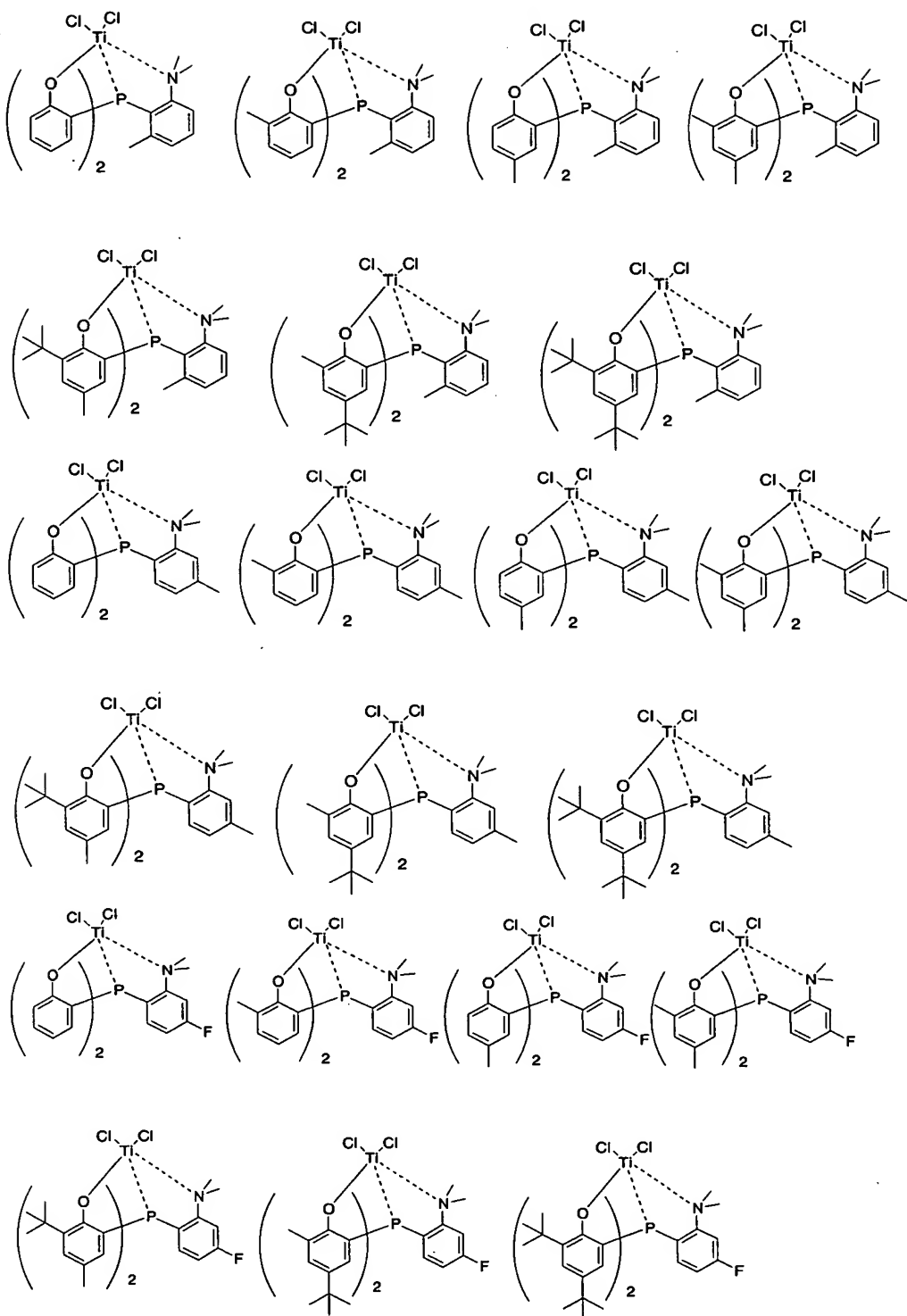


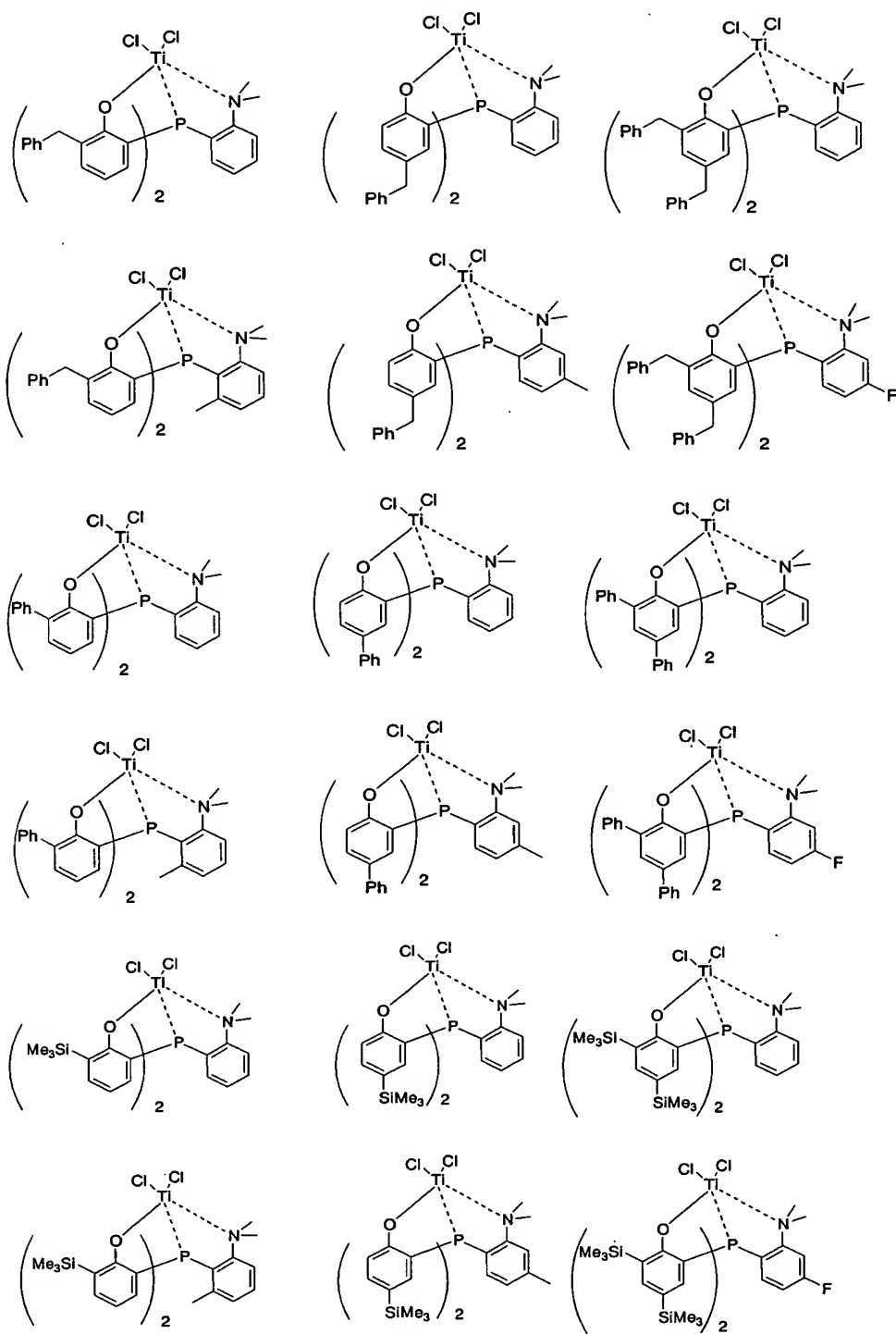




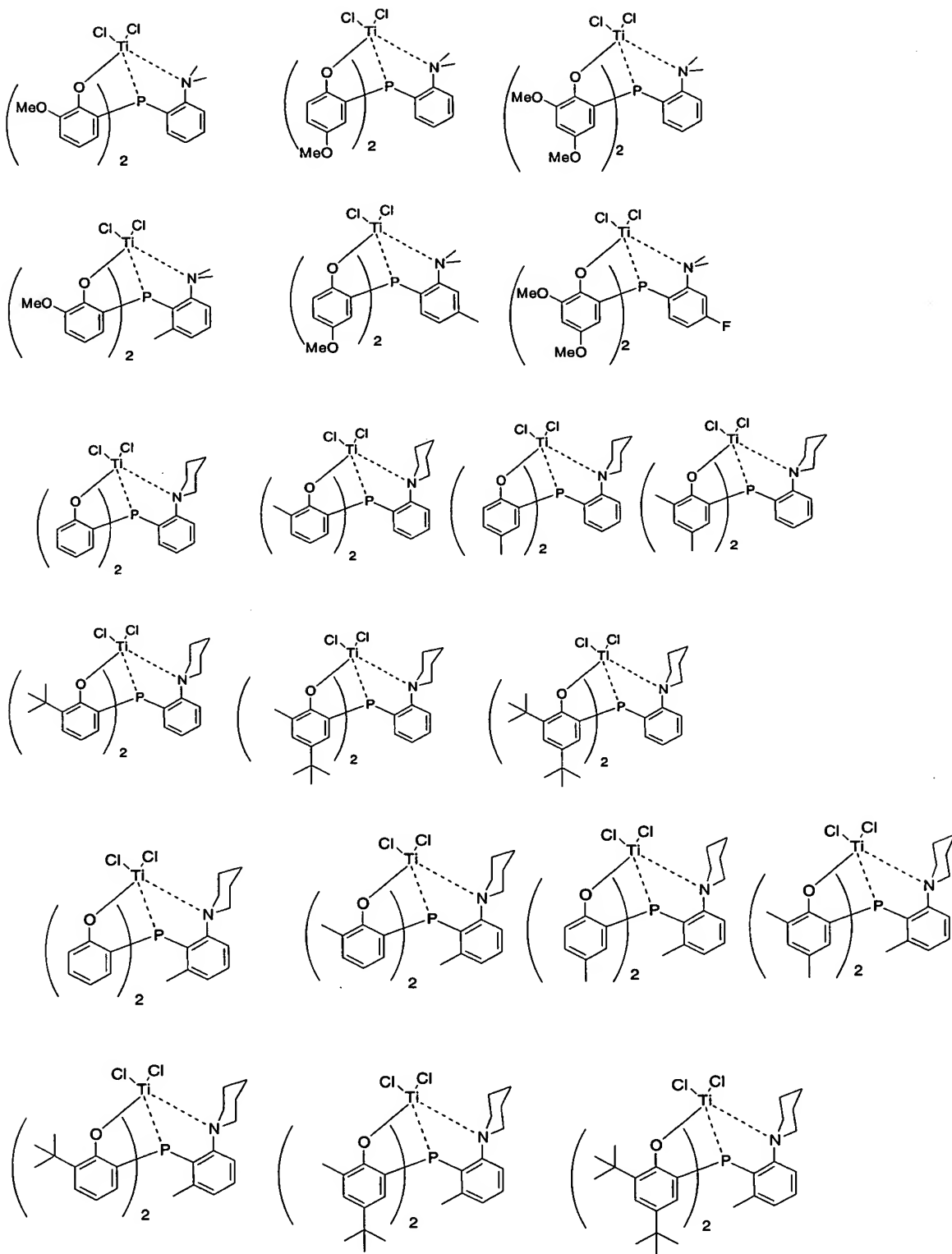


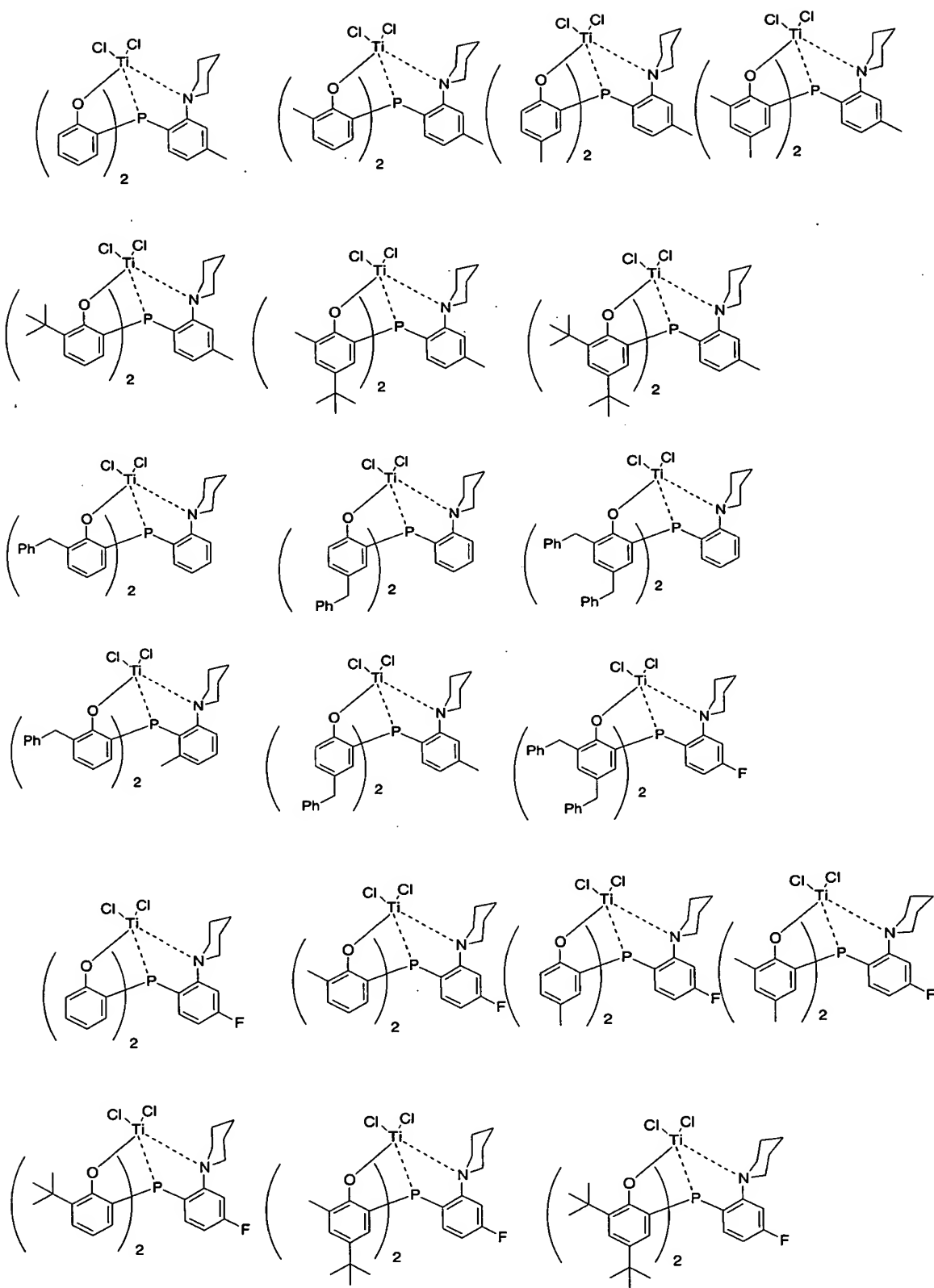


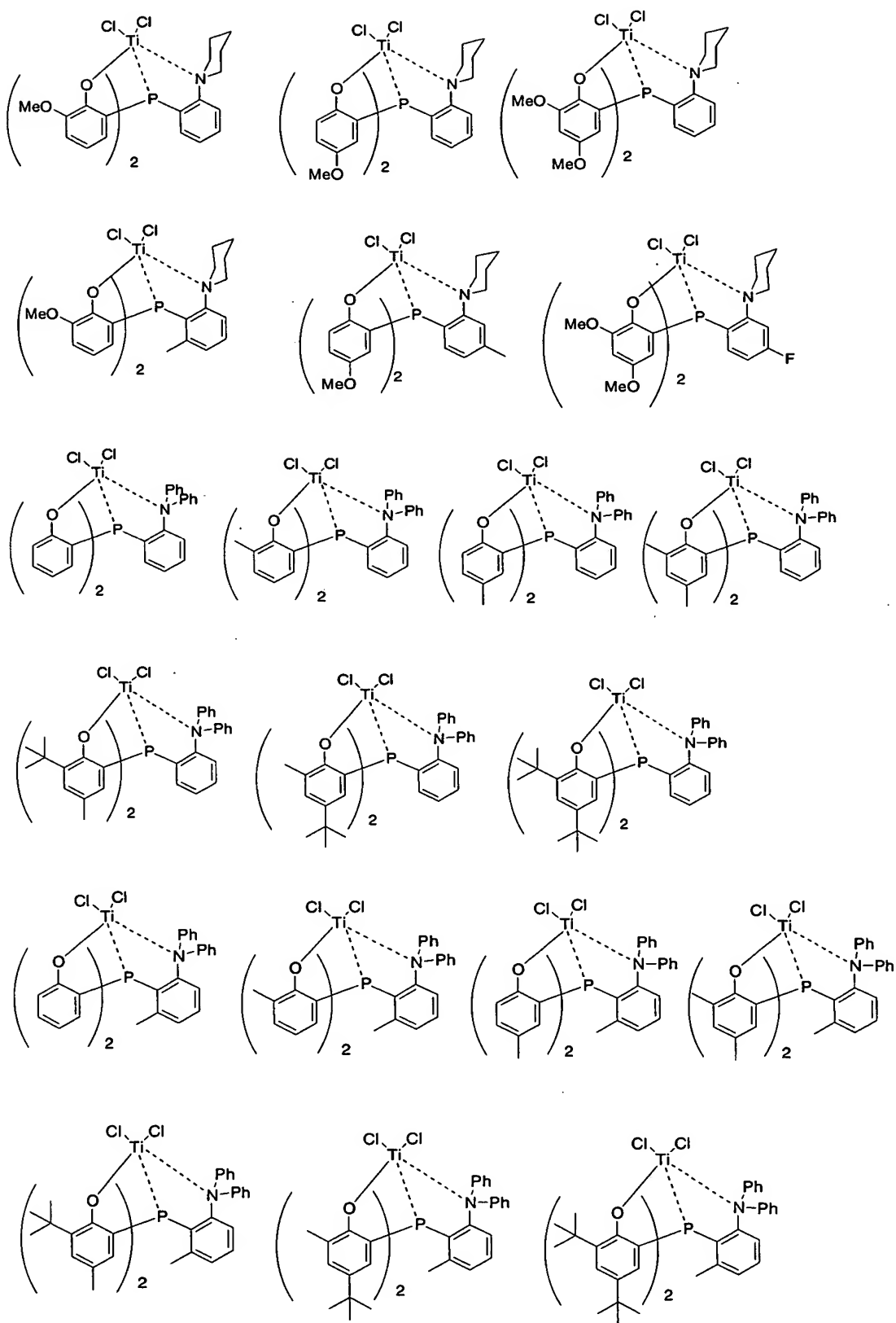


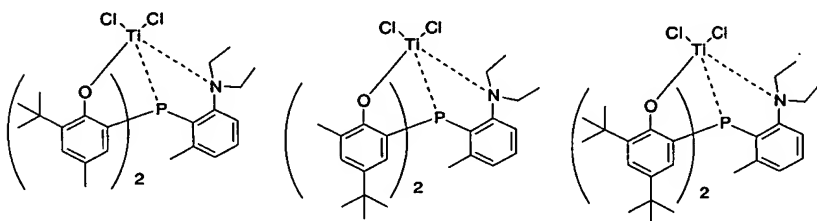
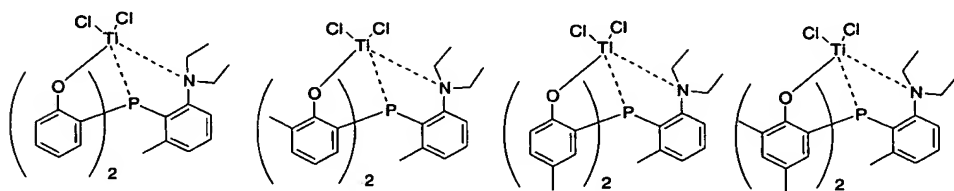
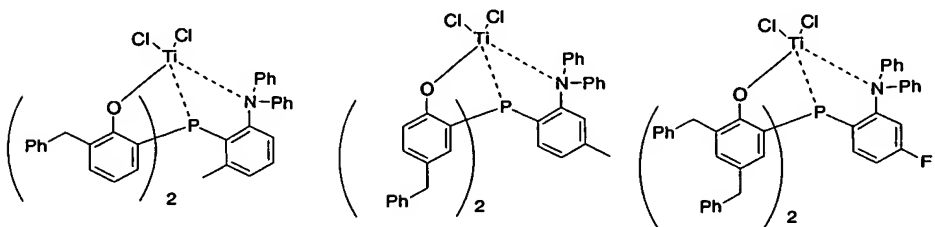
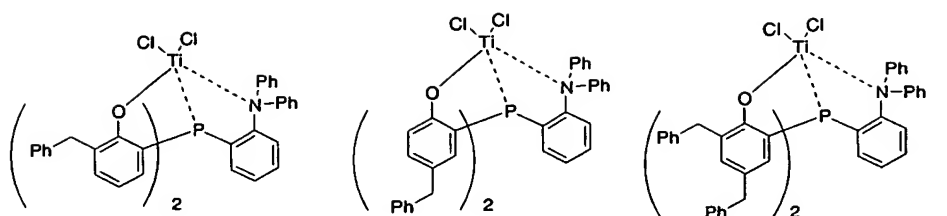


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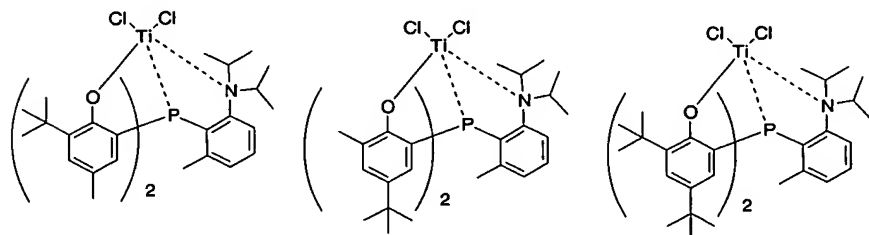
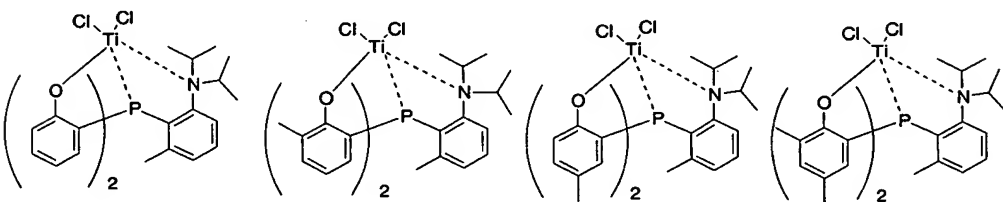


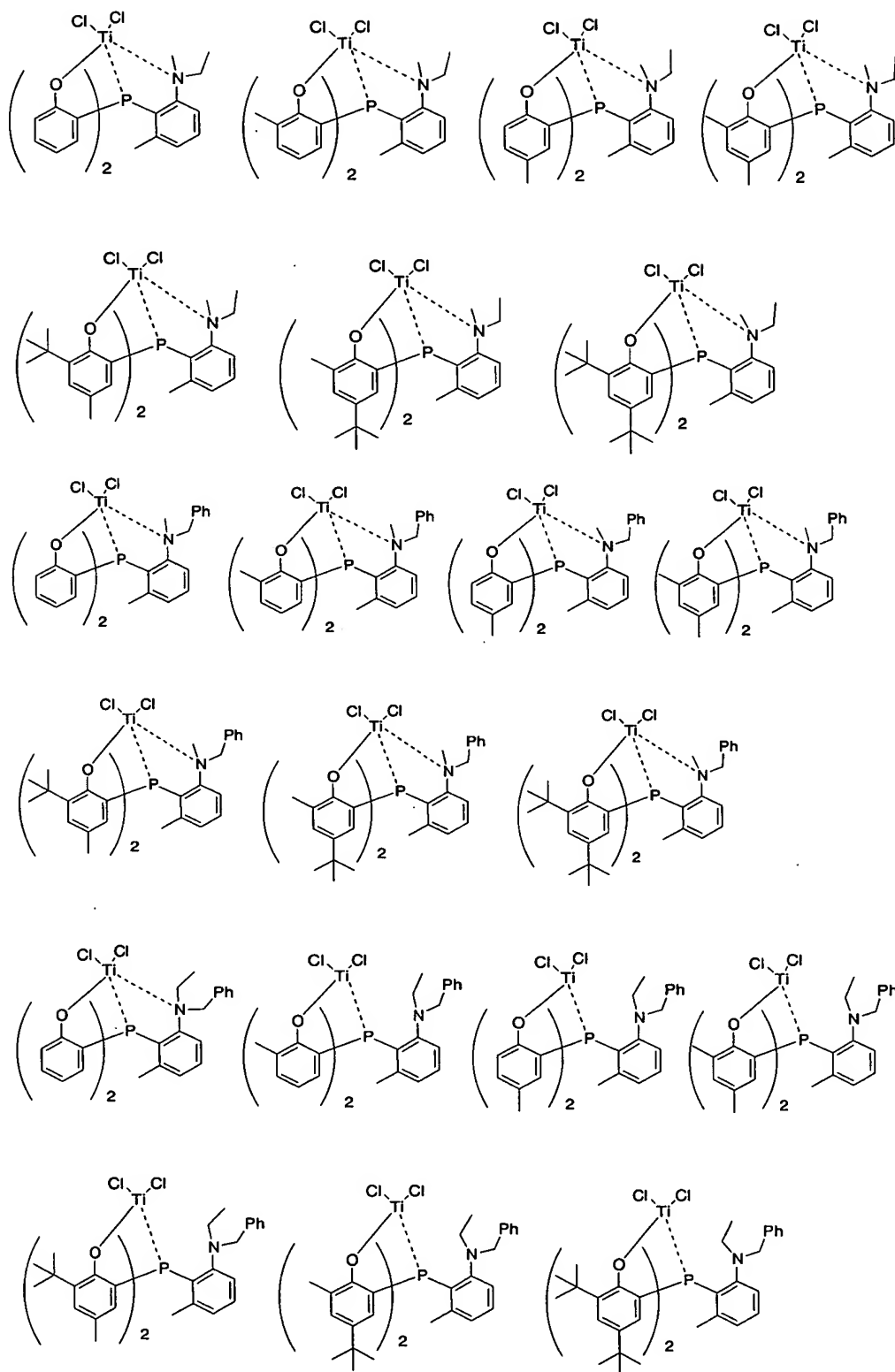


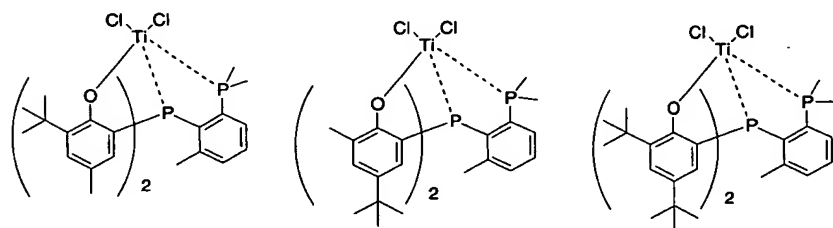
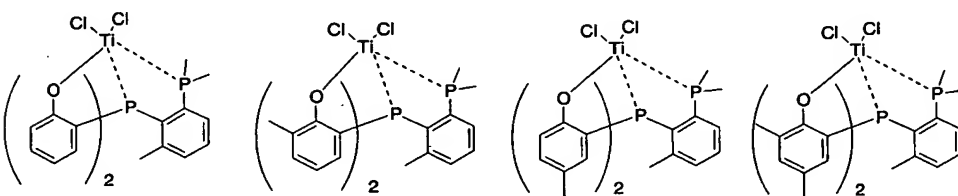
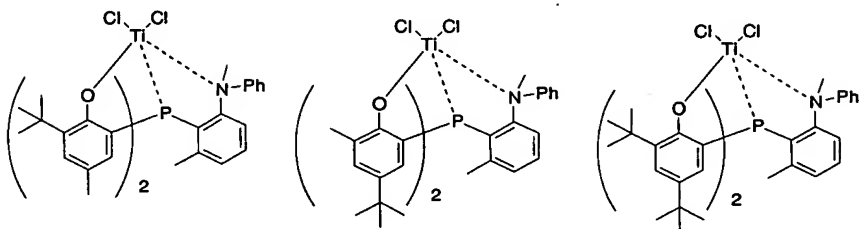
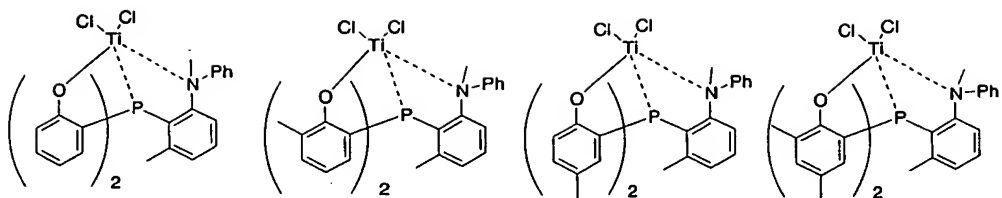


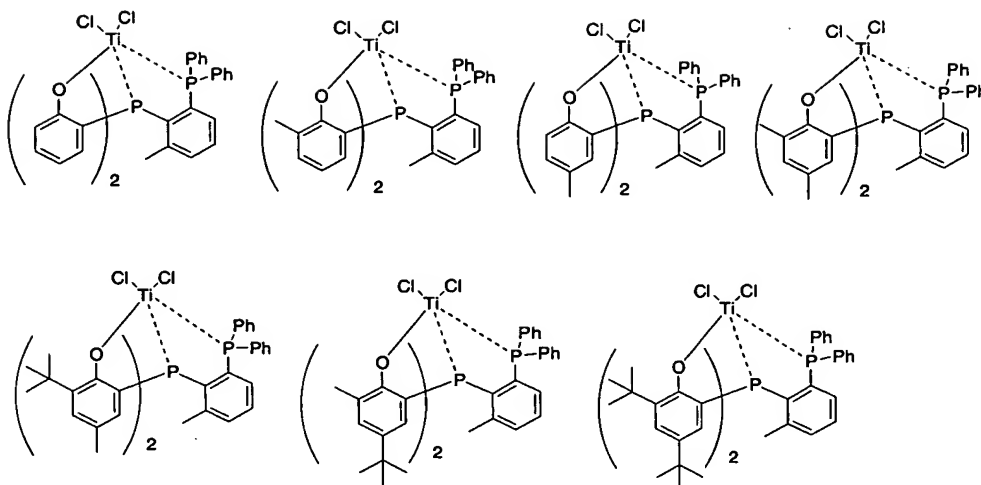


5









The compounds in which the titanium atom is replaced by the zirconium atom or hafnium atom can also be exemplified.

In the polymerization reaction, the transition metal complex thus produced may be charged for use with compound A or compound B as an additional component(s) in an optional order, or a products obtained by contacting the components optionally selected from the components prior to the polymerization.

Organic aluminum compound known in the art may be used as compound A in the invention. Preferably, the organic aluminum compound known in the art may be used as compound A, and any one of compounds A1 to A3, or a mixture of at least two of them is preferable.

Examples of organic aluminum compound A1 represented by E1a-A1-Z3a include trialkylaluminum such as trimethylaluminum, triethylaluminum, tripropylaluminum, triisobutylaluminum, trihexylaluminum or the like; dialkyl aluminum chloride such as dimethylaluminum chloride, diethylaluminum chloride, dipropylaluminum chloride, diisobutylaluminum chloride, dihexylaluminum chloride or the like; alkylaluminum dichloride such as methylaluminum dichloride, ethylaluminum dichloride,

propylaluminum dichloride, isobutylaluminum dichloride, hexylaluminum dichloride or the like; and dialkylaluminum hydride such as dimethylaluminum hydride, diethylaluminum hydride, dipropylaluminum hydride, diisobutylaluminum hydride, dihexylaluminum hydride or the like. Trialkylaluminum is preferable, and triethylaluminum and triisobutylaluminum are more preferable.

Specific examples of E2, E3 in cyclic aluminoxane (A2) having the structure of formula $[-A(E2)-O-]_b$ and in linear aluminosilane ((A3) having the structure of formula $E3[-Al(E3)-O-]_c$, respectively, include alkyl groups such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, n-pentyl, neopentyl or the like. b is an integer of 2 or more, and c is an integer of 1 or more. Preferably, E2 or E3 is a methyl or isobutyl group, and b is 2 to 40 and c is 1 to 40, respectively. Specific examples of aluminosilane include methyl aluminosilane (MAO), modified aluminosilane (MMAO) and butyl aluminosilane (BAO).

The aluminosilane can be produced by various methods. The method is not particularly restricted, and the compound may be produced according to the method known in the art. For example, the compound is by reacting a solution prepared by dissolving trialkylaluminum (for example trimethylaluminum) in a suitable solvent (such as benzene and aliphatic hydrocarbon), with water. In another example, the compound is produced by contacting trialkylaluminum (for example trimethylaluminum) with a metal salt containing crystallization water (for example copper sulfate hydrate).

In the boron compound (B1) of formula $BQ^1Q^2Q^3$, Q^1 to Q^3

are preferably halogen atoms, hydrocarbons having 1 to 20 carbon atom(s), or halogenated hydrocarbons having 1 to 20 carbon atom(s).

Specific examples of B1 include

- 5 tris(pentafluorophenyl)borane,
tris(2,3,5,6-tetrafluorophenyl)borane,
tris(2,3,4,5-tetrafluorophenyl)borane,
tris(3,4,5-trifluorophenyl)borane,
tris(2,3,4-trifluorophenyl)borane and
10 phenylbis(pentafluorophenyl)borane. Preferred is
tris(pentafluorophenyl)borane.

In the boron compound (B2) of formula $Z^+(BQ^1Q^2Q^3Q^4)^-$, examples of Q^1 to Q^4 are the same as those exemplified for Q^1 to Q^3 in the boron compound (B1).

- 15 In the specific example of the compound represented by $Z^+(BQ^1Q^2Q^3Q^4)^-$, examples of inorganic cation Z^+ include ferrocenium cation, alkyl-substituted ferrocenyl cation and silver cation, and examples of organic cation Z^+ include triphenylmethyl cation. Examples of $(BQ^1Q^2Q^3Q^4)^-$ include
20 tetrakis(pentafluorophenyl) borate,
tetrakis(2,3,5,6-tetrafluorophenyl) borate,
tetrakis(2,3,4,5-tetrafluorophenyl)
borate, tetrakis(3,4,5-trifluorophenyl) borate,
tetrakis(2,2,4-trifluorophenyl) borate,
25 phenylbis(pentafluorophenyl) borate and
tetrakis(3,5-bistrifluoromethylphenyl) borate.

Specific examples of the combination thereof include ferrocenium tetrakis(pentafluorophenyl) borate, 1,1'-dimethylferrocenium tetrakis(pentafluorophenyl) borate,

silver tetrakis(pentafluorophenyl) borate,
triphenylmethyltetrakis(pentafluorophenyl) borate, and
triphenylmethyltetrakis(3,5-bistrifluoromethylphenyl)
borate. Triphenylmethyltetrakis(pentafluorophenyl) borate
5 is preferable.

In the boron compound (B3) represented by $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$,
 Q^1 to Q^4 are the same as Q^1 to Q^3 in B1 above.

As the specific examples of the compound represented
by $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$, exemplified are those composed of Brønsted
10 acid $(L-H)^+$ such as trialkyl-substituted ammonium, N,N-dialkyl
anilinium, dialkyl ammonium or triaryl phosphonium, and
 $(BQ^1Q^2Q^3Q^4)^-$ as exemplified above.

Specific examples of the combination thereof include
triethylammonium tetrakis(pentafluorophenyl) borate,
15 tripropylammonium tetrakis(pentafluorophenyl) borate,
tri(n-butyl)ammonium tetrakis(pentafluorophenyl) borate,
tri(n-butyl)ammonium tetrakis(3,5-bistrifluoromethylphenyl)
borate, N,N-dimethylanilinium tetrakis(pentafluorophenyl)
borate, N,N-diethylanilinium tetrakis(pentafluorophenyl)
20 borate, N,N-2,4,6-pentamethylanilinium
tetrakis(pentafluorophenyl) borate, N,N-dimethylanilinium
tetrakis ammonium tetrakis(3,5-bistrifluoromethylphenyl)
borate, diisopropylammonium tetrakis(pentafluorophenyl)
borate, dicyclohexylammonium tetrakis(pentafluorophenyl)
25 borate, triphenylphosphonium tetrakis(pentafluorophenyl)
borate, tri(methylphenyl)phosphonium
tetrakis(pentafluorophenyl) borate,
tri(methylphenyl)phosphonium tetrakis(pentafluorophenyl)
borate, and tri(dimethylphenyl)phosphonium

tetrakis(pentafluorophenyl) borate. Preferred are tri(n-butyl)ammonium tetrakis(pentafluorophenyl) borate, N,N-dimethylanilinium tetrakis(pentafluorophenyl) borate and the like.

5 The ratio of each catalyst component is desirably in the range of 0.1 to 10,000, more preferably in the range of 5 to 2000, for the molar ratio of compound A/transition metal complex (1), and more preferably in the range of 0.01 to 100, preferably in the range of 0.5 to 10, for the molar ratio of compound
10 B/transition metal complex (1).

 The concentration of each catalyst component used as a solution is desirably in the range of 0.0001 to 5 mmol/L, preferably in the range of 0.001 to 1 mmol/L, for transition metal complex (1); in the range of 0.01 to 500 mmol/L, preferably
15 in the range of 0.1 to 100 mmol/L in terms of aluminium, for compound A; and in the range of 0.0001 to 5 mmol/L, preferably in the range of 0.001 to 1 mmol/L, for compound B.

 Any one of olefin and diolefin having 2 to 20 carbon atoms can be used as the monomer for polymerization in the invention,
20 and at least two monomers can be used together. While the monomers are exemplified below, the invention is not restricted to these monomers. Specific examples of olefin include ethylene, propylene, butene-1, pentene-1, hexene-1, heptene-1, octene-1, nonen-1, decen-1, 5-methyl-2-pentene-1, vinyl cyclohexene and
25 the like. Examples of the diolefin compound include conjugated diene and non-conjugated diene of the hydrocarbon compound, and specific examples of the non-conjugated diene compound include 1,5-hexadiene, 1,4-hexadiene, 1,4-pentadiene, 1,7-octadiene, 1,8-nonadiene, 1,9-decadiene, 4-methyl-1,4-hexadiene,

5-methyl-1,4-hexadiene, 7-methyl-1,6-octadiene,
5-ethylidene-2-norbornene, dicyclopentadiene,
5-vinyl-2-norbornene, 5-methyl-2-norbornene, norbornadiene,
5-methylene-2-norbornene, 1,5-cyclooctadiene,
5 5,8-endomethylene hexahydronaphthalene and the like; and
specific examples of the conjugated diene compound include
1,3-butadiene, isoprene, 1,3-hexadiene, 1,3-octadiene,
1,3-cyclooctadiene, 1,3-cyclohexadiene and the like.

Specific examples of the monomer constituting the
10 copolymer include ethylene and propylene, ethylene and butene-1,
ethylene and hexene-1, and propylene and butene-1, as well as
combinations further using 5-ethylidene-2-norbornene thereto,
the invention is not restricted to these compounds.

Aromatic vinyl compounds can be used as the monomer in
15 the invention. Specific examples of the aromatic vinyl compound
include styrene, o-methylstyrene, m-methylstyrene,
p-methylstyrene, o,p-dimethylstyrene, o-ethylstyrene,
m-ethylstyrene, p-ethylstyrene, o-chlorostyrene,
p-chlorostyrene, α -methylstyrene, divinylstyrene and the like.

20 The polymerization method is not particularly restricted,
and the polymerization can be conducted by solvent
polymerization using an aliphatic hydrocarbon such as butane,
pentane, hexane, heptane or octane, aromatic hydrocarbon such
as benzene or toluene, or halogenated hydrocarbon such as
25 methylene dichloride as the solvent, slurry polymerization or
gas phase polymerization using gaseous monomers. Continuous
polymerization or batch-wise polymerization can be used.

The polymerization temperature can be set in the range
of -50°C to 200°C. The temperature range of -20°C to 100°C is

preferable. The polymerization reaction pressure is preferably in the range of atmospheric pressure to 6 MPa (60 kg/cm²G). The polymerization reaction time is appropriately selected depending on the kind of the desired polymer and reaction equipment, and it may be in the range of 1 minute to 20 hours. A chain transfer agent such as hydrogen can be added in the invention for controlling the molecular weight of the copolymer.

Examples

While the invention is described in more detail with reference to examples, the invention is not restricted to these examples. The properties of the polymers in the examples were measured by the following methods.

[Molecular weight and molecular weight distribution]

The molecular weight and molecular weight distribution were measured as follows using Rapid GPC (trade name; manufactured by Symyx Co.).

Pump: (LC pump), manufactured by Gilson Co.

Model 1305 (trade name), pump head 25.SC

Column: PLgel Mixed-B (trade name; manufactured by Polymer Laboratories (PL) Co.), 10 μ m,

7.5 mm ϕ \times 300 mm

Mobile phase: o-dichlorobenzene

Dissolving solvent: 1,2,4-trichlorobenzene

Flow rate: 2 ml/min

Column temperature: 160°C

Calibration curve: polystyrene (PS, standard manufactured by PL Co.), 8 samples

Standard molecular weight of PS; 5,000, 10,050, 28,500,

65,500, 185,400, 483,000, 1,013,000, 3,390,000

[Melting point]

Melting point was measured under the following condition
using SAMMS (Sensor Array Modular System, trade name,
5 manufactured by Symyx Co.)

Measurement mode: melting temperature measurement by heat
capacity spectroscopy

Atmospheric gas: vacuum (3.0×10^{-4} Torr or below)

Temperature program:

10 (start) room temperature

(rate of temperature increase) about 50°C/min

(hold) 200°C (0 minute)

[Me branching]

Me branching was measured under the following condition
15 using IR spectrometer (EQINOX 55, trade name, manufactured by
Bruker Co.)

Measurement mode: reflection-transmission method (a film
is formed on a mirror surface)

Blank: mirror surface (air)

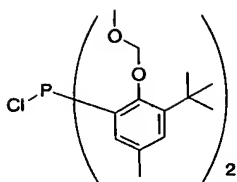
20 Measuring condition:

(resolution) 2 cm^{-1}

(number of integration) 128 times

(wavelength) 400 to 4000 cm^{-1}

25 Example A1: Synthesis of compound A1



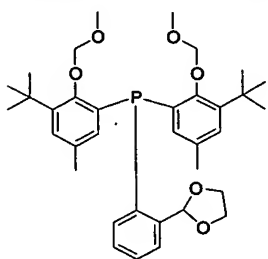
A1

A 1.56 M hexane solution (7.05 mL) of n-butyl lithium was added dropwise into a tetrahydrofuran solution (23.5 mL) of 1-methoxymethoxy-2-tert-butyl-4-methylbenzene (2.08 g, 10.0 mmol) at -78°C , and the solution was warmed to room temperature with stirring for 1 hour. The reaction solution was added into a tetrahydrofuran solution (23.5 mL) of phosphorous trichloride (0.69 g, 5.0 mmol) at -78°C , and the solution was warmed to room temperature with stirring for 5 hours. Compound A1 was quantitatively obtained by removing the solvent under a reduced pressure after removing insoluble substances by filtration.

^1H NMR(CD_2Cl_2) δ 1.38(18H), 2.25(6H), 3.60(6H), 5.06-5.26(4H), 7.07-7.27(4H)

^{31}P NMR(CD_2Cl_2) 79.15

Example A2: Synthesis of compound A2



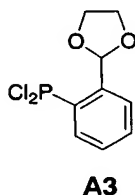
A2

A 1.56 M hexane solution (35.3 mL) of n-butyl lithium was added dropwise into a tetrahydrofuran solution (180.6 mL) of 2-(o-bromophenyl)-1,3-dioxolane (11.15 g, 50.0 mol) at -78°C ,

and the solution was warmed to room temperature with stirring for 2 hours. The reaction mixture was cooled to -78°C , and a tetrahydrofuran solution (77.4 mL) of compound A1 (24.05 g, 50.0 mmol) was added therein followed by warming to room temperature with stirring for 10 hours. The reaction was stopped by adding deionized water (200.0 mL) and toluene (200.0 mL), and the solvent was removed after washing the organic layer with saturated aqueous sodium chloride solution (100 mL) to obtain a crude product as a pale yellow oil. The crude product was purified by silica gel column chromatography (hexane/ethyl acetate = 30/1 to 4/1) to obtain compound A2 (9.52 g, yield 32.0%) as a white solid.

^1H NMR(CDCl_3) δ 1.37(18H), 2.10(6H), 3.45(6H), 3.93-4.14(4H), 5.11-5.13(4H), 5.20(1H), 6.34(2H), 6.44(2H), 6.92-7.66(4H)

Example A3: Synthesis of compound A3



A 1.56 M hexane solution (21.2 mL) of n-butyl lithium was added dropwise into a diethylether solution (145.0 mL) of 2-(o-bromophenyl)-1,3-dioxane (6.87 g, 30.0 mmol) at -78°C , and the solution was warmed to room temperature with stirring for 2 hours. The mixed reaction solution was cooled to -78°C , and a diethylether solution (116.0 mL) of phosphorous trichloride (8.24 g, 60.0 mmol) was added to the solution followed by warming to room temperature with stirring for 10 hours. Compound A3

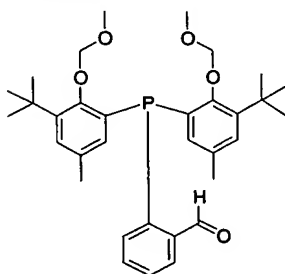
was obtained by removing the solvent by evaporation in vacuum after removing insoluble materials by filtration.

^{31}P (CD_2Cl_2): δ 160.8

5 Example A4: Synthesis of compound A2

A 1.56 M hexane solution (33.8 mL) of n-butyl lithium was added dropwise into a tetrahydrofuran solution (158 mL) of 2-tert-butyl-1-methoxymethoxy-4-methylbenzene (10.0 g, 48.0 mmol) at -78°C , and the solution was warmed to room temperature with stirring for 2 hours. The reaction mixture was cooled to -78°C , and a tetrahydrofuran solution (67.5 mL) of compound A3 (6.03 g, 24.0 mmol) was added into the cooled solution, followed by warming to room temperature with stirring for 10 hours. Compound A2 was obtained by applying the same post treatment to the resulting mixture as in Example A2.

Example A5: Synthesis of compound A4



A4

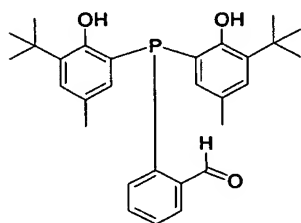
Into a solution of compound A2 (1.49 g, 2.50 mmol) in a mixed solvent of tetrahydrofuran/water = 10/1 (46.3 mL), 98% sulfuric acid (1.32 g) was added at room temperature and the mixture was stirred for 3 hours. The reaction was stopped by adding deionized water (70.0 mL) and toluene (50.0 mL). After

washing the organic layer with saturated aqueous sodium chloride solution (70 mL), the solvent was removed by evaporation to quantitatively obtain compound A4 as a pale yellow oil.

$^1\text{H NMR}(\text{CDCl}_3)$ δ 1.40(18H), 2.10(6H), 3.50(6H), 5.26(4H), 6.27(2H), 6.97-7.98(6H),

5 10.6(1H)

Examples A6: Synthesis of compound A5



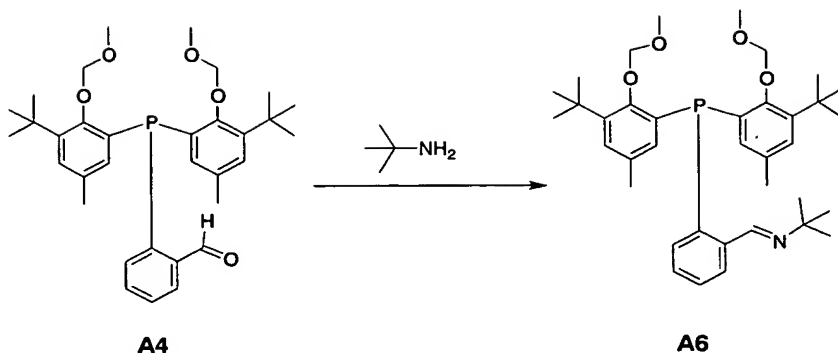
A5

Acetyl chloride (1.96 g, 25.0 mmol) was added to a solution
10 of compound A4 (2.75 g, 5.00 mmol) in a mixed solvent of ethyl acetate and methanol (1:1, 110.0 mL), and then the mixture was stirred for 15 hours at room temperature. A crude product was obtained as an yellow oil by removing the solvent by evaporation in vacuum. The crude product was purified by silica gel column
15 chromatography to obtain 0.64 g of compound A5 (yield 64.0%) as an yellow solid.

$^1\text{H NMR}(\text{CDCl}_3)$ δ 1.34(18H), 2.09(6H), 6.33(2H), 6.51(2H), 7.07-7.87(6H), 10.1(1H)

$^{31}\text{P NMR}(\text{C}_6\text{D}_6)$ δ -52.8

20 Example A7: Synthesis of compound A6

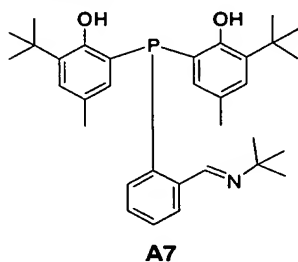


Tert-butylamine (0.91 g, 12.5 mmol) was added to a solution of compound A4 (1.38 g, 2.50 mmol) in ethanol solution (62.4 mL), and the mixture was heated to 40°C and then stirred for 5 hours. Compound A6 was quantitatively obtained by removing

¹H NMR(CDCl₃) δ 1.15(9H), 1.40(18H), 2.12(6H), 3.50(6H), 5.10-5.19(4H), 6.47(2H), 6.90-7.95(6H), 8.90(1H)

¹³C NMR(CDCl₃) δ 21.1, 29.5, 30.8, 36.1, 57.2, 57.6, 99.5, 126.4, 142.8, 154.5, 156.3

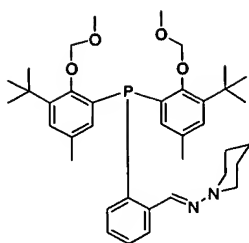
10 Example A8: Synthesis of compound A7



Compound A7 can be obtained by adding acetyl chloride to a solution of compound A6 in a mixed solution of ethyl acetate and methanol (1:1) with stirring at room temperature, followed by removing the solvent under a reduced pressure.

15

Example A9: Synthesis of compound A8

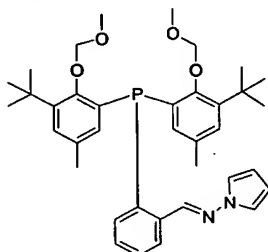


A8

Compound A8 was quantitatively obtained by the same manner as in Example A7, except that aminopiperidine was used in place of tert-butylamine.

¹H NMR(CDCl₃) δ 1.38-3.03(10H), 1.40(18H), 2.12(6H), 3.48(6H), 5.08-5.18(4H), 6.50(2H),
 5 6.89-7.92(6H), 8.11(1H)

Example A10: Synthesis of compound A9

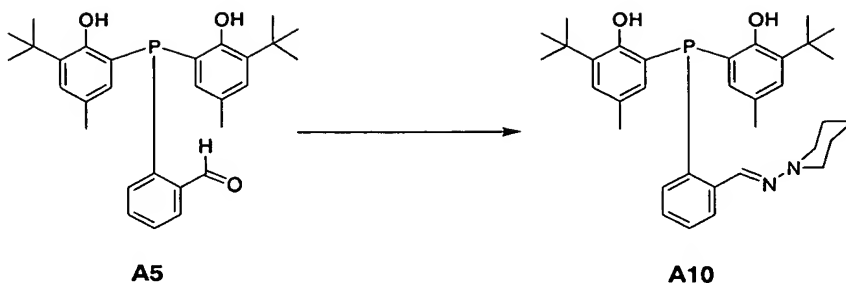


A9

Compound A9 was quantitatively obtained by the same manner as in Example A7, except that aminopyrrole was used in place
 10 of tert-butylamine.

¹H NMR(CDCl₃) δ 1.40(18H), 2.13(6H), 3.51(6H), 5.09-5.24(4H), 6.48(2H), 7.01-8.12(10H),
 9.15(1H)

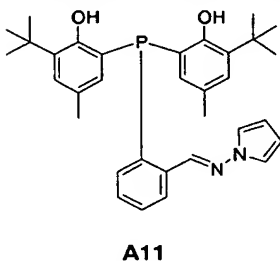
15 Example A11: Synthesis of compound A10



Aminopiperidine (0.03 g, 0.25 mmol) was added to a solution of compound A5 (0.12 g, 0.25 mmol) in ethanol solution (44.0 mL) at 0°C with stirring for 3 hours. Compound A10 was quantitatively obtained by removing the solvent by evaporation.

5

Example A12: Synthesis of compound A11



Compound A11 was quantitatively obtained by the same manner as in Example A11, except that aminopyrrole was used in place of aminopiperidine.

10 $^1\text{H NMR}(\text{CDCl}_3)$ δ 1.41(18H), 2.16(6H), 6.15(2H), 6.67(2H), 6.96-7.49(10H), 8.68(1H)

Example A13: Synthesis of compound A10

Compound A10 can be obtained by adding acetyl chloride to a solution of compound A8 in a mixed solvent of ethyl acetate and methanol (1:1) at room temperature with stirring, and by removing the solvent by evaporation in vacuum.

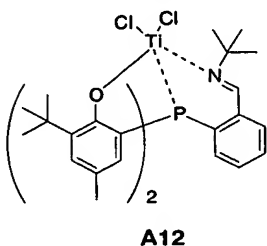
15

Example A14: Synthesis of compound A11

Compound A11 can be obtained by adding acetyl chloride to a solution of compound A9 in a mixed solvent of ethyl acetate and methanol (1:1) at room temperature with stirring, and by removing the solvent by evaporation in vacuum.

5

Example A15: Synthesis of complex A12

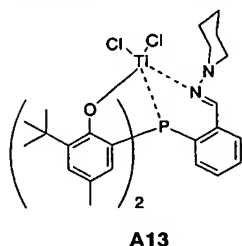


A toluene solution (2.31 mL) of titanium tetrachloride (0.08 g, 0.40 mmol) was added dropwise into a toluene solution (2.31 mL) of compound A6 (0.20 g, 0.33 mmol) at -78°C , and the solution was stirred for 10 hours after warming to room temperature. Compound 12 (204.7 mg, yield 97.5%) was obtained as a red solid by removing the solvent under a reduced pressure after removing insoluble substances by filtration.

^{31}P NMR(C_6D_6) δ 22.8

15 EI-MS 635(M^{+1})

Example A16: Synthesis of complex A13



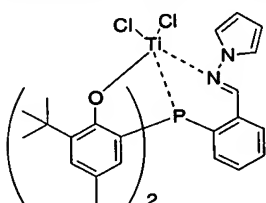
Complex A13 (289.3 mg) was obtained in 87.8% yield by the

same manner as in Example A15, except that compound A8 was used in place of compound A6.

^1H NMR(CD_2Cl_2) δ 1.27-2.00(10H), 1.38(18H), 2.34(6H), 6.86(2H), 7.06(2H), 7.48-8.15(4H), 10.23(1H)

5 ^{31}P NMR(C_6D_6) δ 7.16
EI-MS 626(M-Cl)

Example A17: Synthesis of complex A14



A14

A 1.57 M solution of n-butyl lithium in hexane (0.64 mL) was added dropwise into a tetrahydrofuran solution (4.45 mL) of compound A11 (0.26 g, 0.50 mmol) at -78°C , and the solution was stirred for 1 hour after warming to room temperature. Then, a tetrahydrofuran solution (4.45 mL) of titanium tetrachloride/2-tetrahydrofuran complex (0.17 g, 0.50 mmol) was dripped into the mixed solution above at -78°C in 2 hours. After warming to room temperature, the reaction mixture was stirred for 10 hours. After removal of the solvent by evaporation in vacuum, toluene (10.0 mL) was added to the residue. After removal of the insoluble materials by filtration, evaporation of the solvent gave complex A14 as a red solid (183.9 mg, yield 57.5%).

^1H NMR(C_6D_6) δ 1.25(9H), 1.47(9H), 1.69(3H), 1.79(3H), 6.29(2H), 6.95-8.42(10H), 9.07(1H)

^{31}P NMR(C_6D_6) δ 9.60

ESI-MS(solvent: CH_3CN) 617(M^+ CH_3CN -pyrolyle)

Examples of polymerization reaction

Example A18:

Toluene (5.0 mL) was added to an autoclave under nitrogen.
5 After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μ mol) and complex A12 (0.10 μ mol) were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer was produced at a rate of 6.4×10^6 g per hour per 1 mole of titanium
10 by the polymerization reaction.

Example A19:

A polymer was produced by the same method as in Example A18, except that a hexane solution of triisobutyl aluminum (40
15 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenyl borane (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 1.1×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

20 Example A20:

A polymer was produced by the same method as in Example A18, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30
25 μ mol) were used in place of MMAO. The polymer was produced at a rate of 3.4×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A21:

A polymer was produced by the same method as in Example A18, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 4.4×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A22:

10 Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex A12 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 20
15 minutes. The polymer with a molecular weight (M_w) of 1.26×10^6 , molecular weight distribution (M_w/M_n) of 4.4 and melting point (T_m) of 126.7°C was produced at a rate of 4.8×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

20 Example A23:

A polymer was produced by the same method as in Example A22, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO.
25 The polymer was produced at a rate of 7.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example A24:

A polymer was produced by the same method as in Example

A22, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 1.97×10^7 , a melting point (T_m) of 124.7°C and a number of branches of Me of 2 per 1,000 carbon atoms was produced at a rate of 2.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

10 Example A25:

A polymer was produced by the same method as in Example A22, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 2.64×10^6 , molecular weight distribution (M_w/M_n) of 1.4, a melting point (T_m) of 121.4°C and a number of branches of Me of 1 per 1,000 carbon atoms was produced at a rate of 3.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

20

Example A26:

Toluene (5.0 mL) and 1-hexene (50 μL) were added to an autoclave under nitrogen. After stabilizing at 70°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmmol) and complex A12 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize. The polymer with a molecular weight (M_w) of 1.42×10^6 , molecular weight distribution (M_w/M_n) of 4.1, melting point (T_m) of 126.0°C and a number of branches of Me of 7 per 1,000 carbon atoms was

produced at a rate of 2.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A27:

5 A polymer was produced by the same method as in Example A26, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer with a molecular weight (Mw) of 2.29×10^6 , molecular weight distribution (Mw/Mn) of 2.2, a melting point (Tm) of 128.2°C and a number of branches of Me of 2 per 1,000 carbon atoms was produced at a rate of 1.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

15 Example A28:

 A polymer was produced by the same method as in Example A26, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer with a molecular weight (Mw) of 1.26×10^6 , molecular weight distribution (Mw/Mn) of 1.5, a melting point (Tm) of 129.8°C and a number of branches of Me of 8 per 1,000 carbon atoms was produced at a rate of 1.5×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example A29:

 Toluene (5.0 mL) and 1-hexene (40 μ L) were added to an autoclave under nitrogen. After stabilizing at 130°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa.

MMAO (100 μmol) and complex A12 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize. The polymer was produced at a rate of 1.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example A30:

A polymer was produced by the same method as in Example A29, except that a hexane solution of triisobutyl aluminum (4 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
10 dimethylanilinium tetrakis(pentafluorophenyl)borate (0.030 μmol) were used in place of MMAO. The polymer was produced at a rate of 8.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

15 Example A31:

A polymer was produced by the same method as in Example A29, except that a hexane solution of triisobutyl aluminum (4 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol)
20 were used in place of MMAO. The polymer was produced at a rate of 8.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example A32:

25 Toluene (5.0 mL) was added to an autoclave under nitrogen and, after stabilizing at 40°C, ethylene was added with compression and was stabilized at 0.60 MPa. MMAO (100 μmol) and complex A13 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize. The polymer was produced

at a rate of 2.7×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A33:

5 A polymer was produced by the same method as in Example A32, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 3.0×10^5 g per hour per
10 1 mole of titanium by the polymerization reaction.

Example A34:

 A polymer was produced by the same method as in Example A32, except that a hexane solution of triisobutyl aluminum (40
15 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 2.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

20

Example A35:

 A polymer was produced by the same method as in Example A32, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and
25 triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 2.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A36:

Toluene (5.0 mL) and 1-hexene (60 μ L) were added to an autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μ mmol) and complex A13 (0.10 μ mol) were added to the autoclave, and the mixture was allowed to polymerize. The polymer with a molecular weight (M_w) of 2.5×10^6 , molecular weight distribution (M_w/M_n) of 25.6 and a melting point (T_m) of 102.0°C was produced at a rate of 2.2×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A37:

A polymer was produced by the same method as in Example A36, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenyl borane (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 3.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example A38:

A polymer was produced by the same method as in Example A36, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 1.7×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A39:

A polymer was produced by the same method as in Example A36, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (Mw) of 2.89×10^6 , molecular weight distribution (Mw/Mn) of 8.4 and a melting point (Tm) of 107.1°C was produced at a rate of 1.9×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A40:

Toluene (5.0 mL) was added to an autoclave under. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmmol) and complex A14 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer was produced at a rate of 4.5×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A41:

A polymer was produced by the same method as in Example A40, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 4.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example A42:

A polymer was produced by the same method as in Example

A40, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at
5 a rate of 5.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A43:

A polymer was produced by the same method as in Example
10 A40, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate
15 of 6.2×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A44:

Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen and, after stabilizing at 40°C , ethylene
20 was added with compression and was stabilized at 0.60 MPa. MMAO (100 μmol) and complex A14 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize. The polymer with a molecular weight (M_w) of 1.94×10^6 , molecular weight
distribution (M_w/M_n) of 2.4 and a melting point (T_m) of 123.0°C
25 was produced at a rate of 1.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A45:

A polymer was produced by the same method as in Example

A44, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 2.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

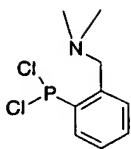
Example A46:

A polymer was produced by the same method as in Example A44, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 4.50×10^6 , molecular weight distribution (M_w/M_n) of 1.3 and a melting point (T_m) of 119.8°C was produced at a rate of 4.2×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example A47:

A polymer was produced by the same method as in Example A44, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 4.33×10^6 , molecular weight distribution (M_w/M_n) of 1.4 and a melting point (T_m) of 127.1°C was produced at a rate of 4.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example B1: Synthesis of compound B1



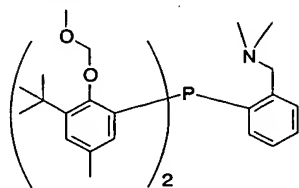
B1

A 1.56 M solution of n-butyl lithium in hexane (14.1 mL) was dripped into an ether solution (57.0 mL) of N,N-dimethylbenzylamine (2.70 g, 20.0 mmol) at 0°C, and the reaction mixture was warmed to room temperature then stirred
 5 for 24 hours. The mixture was cooled to -78°C, and an ether solution (77.0 mL) of phosphorous trichloride (5.49 g, 40.0 mmol) was added followed by warming to room temperature then stirred for 2 hours. Compound B1 was quantitatively obtained by removing the solvent from the filtrate in vacuum after removing insoluble
 10 materials by filtration.

$^1\text{H NMR}(\text{CD}_2\text{Cl}_2)$ δ 2.46(6H), 4.02(2H), 7.29-8.53(4H)

$^{31}\text{P NMR}(\text{CD}_2\text{Cl}_2)$ 115.6

Examples B2: Synthesis of compound B2



B2

15 A 1.56 M of n-butyl lithium solution in hexane (28.2 mL) was added dropwise into a tetrahydrofuran solution (131.4 mL) of 2-tert-butyl-1-methoxymethoxy-4-methylbenzne (8.33 g, 40 mmol) at -78C, and the mixture was warmed to room temperature then stirred for 1 hour. The reaction mixture was cooled to
 20 -78°C, and a tetrahydrofuran solution (56.3 mL) of compound B1

(4.72 g, 20.0 mmol) was added followed by warming to room temperature with stirring for 10 hours. The reaction was stopped by adding deionized water (100.0 mL) and toluene (100 mL). The organic layer was washed with saturated aqueous sodium chloride solution (100 mL) followed by removing the solvent by evaporation to obtain a pale yellow oil as a desired product. The product was purified by silica gel column chromatography (hexane/ethyl acetate = 10/1) to obtain compound B2 as a white solid (4.35 g, yield 37.5%).

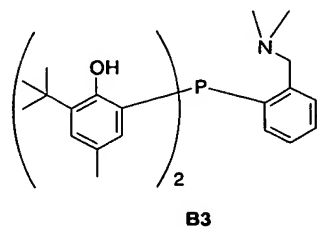
$^1\text{H NMR}(\text{CDCl}_3)$ δ 1.40(18H), 2.09(6H), 2.10(6H), 3.50(6H), 3.55(2H), 5.16-5.19(4H), 6.35(2H), 6.86-7.47(6H)

MS 536(M+1)

Example B3: Synthesis of compound B2

A hexane solution (1.56 M) of n-butyl lithium was dripped into an ether solution of N,N-dimethylbenzylamine at 0°C, and the solution is warmed to room temperature with stirring for 24 hours. The mixture is cooled to -78°C, and an ether solution of compound A1 is added followed by warming to room temperature with stirring for 10 hours. Compound B2 is obtained by removing the solvent from the filtrate in vacuum after removing insoluble substances by filtration.

Example B4: Synthesis of compound B3



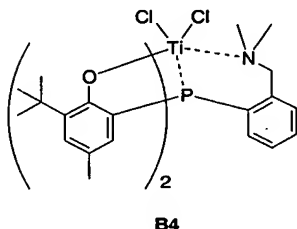
Acetyl chloride (0.79 g, 10.0 mmol) was added to a solution of compound B2 (0.95 g, 1.64 mol) in a mixed solvent (57.0 mL) of ethyl acetate and methanol (1/1) at room temperature, and the solution was stirred at room temperature for 15 hours.

- 5 Compound B3 was obtained as a white solid (403,5 mg, yield 49.8%) by removing the solvent by evaporation in vacuum.

$^1\text{H NMR}(\text{CDCl}_3)$ d1.41(18H), 2.25(6H), 3.01(6H), 4.63(2H), 6.31(2H), 7.06-8.81(6H)

$^{31}\text{P NMR}(\text{C}_6\text{D}_6)$ d-26.9

10 **Example B5: Synthesis of complex B4**



A toluene solution (6.70 mL) of titanium tetrachloride (0.40 g, 2.10 mmol) was added dropwise into a toluene solution (6.70 mL) of compound B2 (0.58 g, 1.00 mmol) at -78°C , and the mixture was stirred at room temperature for 10 hours. Complex

- 15 B4 was quantitatively obtained as a brown solid by washing with pentane (2 mL) after removing the solvent by distillation.

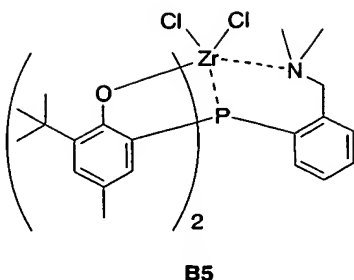
$^1\text{H NMR}(\text{C}_6\text{D}_6)$ d1.59(18H), 2.05(6H), 2.41(6H), 3.77(2H), 6.84(2H), 6.99-7.89(6H)

$^{31}\text{P NMR}(\text{C}_6\text{D}_6)$ d28.3

EI-MS 607(M-1)

20

Example B6: Synthesis of complex B5



A n-butyl lithium solution (1.56 M) in hexane (1.53 mL) was added into a solution of compound B3 (0.42 g, 0.80 mmol) in tetrahydrofuran (7.14 mL) at -78°C , and the mixture was warmed to room temperature with stirring for 1 hour. The reaction mixture was added into a solution of zirconium tetrachloride bis(tetrahydrofuran) complex (0.30 g, 0.80 mol) in tetrahydrofuran (7.14 mL) at -78°C . After stirring the solution for 10 hours at room temperature, 10.0 mL of toluene was added. Complex B5 was obtained as a white solid (270 mg, yield 50%) by removing the solvent from the filtrate in vacuum after removing insoluble materials by filtration.

EI-MS: 649 (M-1)

Examples of polymerization reaction

Example B7:

Toluene (5.0 mL) was added in an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. Methylaluminoxane (100 μmol) and complex B4 (0.10 μmol) were added in the autoclave to polymerize the mixture for 5 minutes. A polymer was produced at a rate of 3.62×10^7 g per hour per 1 mol of titanium by the polymerization reaction.

Example B8:

A polymer was produced by the same method as in Example B6 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 6.00×10^5 g per hour per 1 mol of titanium by polymerization.

10 Example B9:

A polymer was produced by the same method as in Example B6 by polymerization for 18 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 6.90×10^6 g per hour per 1 mol of titanium by polymerization.

Example B10:

20 A polymer was produced by the same method as in Example B6 by polymerization for 14 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 8.80×10^6 g per hour per 1 mol of titanium by polymerization.

Example B11:

Toluene (5.0 mL) and 1-hexene (60 μL) were added in an

autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. Methyl aluminoxane (100 μmol) and complex B4 (0.10 μmol) were added in the autoclave to polymerize the mixture for 9 minutes.

5 A polymer was produced at a rate of 2.01×10^7 g per hour per 1 mol of titanium by the polymerization reaction.

Example B12:

A polymer was produced by the same method as in Example B10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 5.00×10^5 g per hour per 1 mol of titanium

15 by polymerization.

Example B13:

A polymer was produced by the same method as in Example B10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of methyl aluminoxane. The polymer with a molecular weight (Mw) of 1.97×10^6 , molecular weight distribution (Mw/Mn) of 1.6

25 and a melting point (T_m) of 117.9°C was produced at a rate of 5.50×10^6 g per hour per 1 mol of titanium by polymerization.

Example B14:

A polymer was produced by the same method as in Example

B10 by polymerization for 14 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of methylaluminoxane. The polymer with a molecular weight (Mw) of 7.61×10^6 , molecular weight distribution (Mw/Mn) of 1.6 and a melting point (T_m) of 113.1°C was produced at a rate of 1.04×10^7 g per hour per 1 mol of titanium by polymerization.

10 Example B15:

Toluene (5.0 mL) was added in an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. Methylaluminoxane (100 μmol) and complex B5 (0.10 μmol) were added in the autoclave to polymerize the mixture for 20 minutes. A polymer was produced at a rate of 4.90×10^6 g per hour per 1 mol of titanium by the polymerization reaction.

Example B16:

20 A polymer was produced by the same method as in Example B14, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of methylaluminoxane. The polymer was produced at a rate of 3.00×10^5 g per hour per 1 mol of zirconium by polymerization.

Example B17:

A polymer was produced by the same method as in Example B14 by polymerization for 20 minutes, except that a hexane

solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 5.50×10^6 g per hour per 1 mol of zirconium by polymerization.

Example B18:

A polymer was produced by the same method as in Example B14 by polymerization for 17 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 7.30×10^6 g per hour per 1 mol of zirconium by polymerization.

Example B19:

Toluene (5.0 mL) and 1-hexene (60 μ L) were added in an autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. Methyl aluminoxane (100 μ mol) and complex B5 (0.10 μ mmol) were added in the autoclave to polymerize the mixture for 20 minutes. A polymer with a molecular weight (M_w) of 2.40×10^5 , molecular weight distribution (M_w/M_n) of 31.2 and a melting point (T_m) of 130.6°C was produced at a rate of 4.60×10^6 g per hour per 1 mol of zirconium by the polymerization reaction.

Example B20:

A polymer was produced by the same method as in Example B18 by polymerization for 20 minutes, except that a hexane

solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μ mol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 4.00×10^5 g per hour per 1 mol of zirconium by polymerization.

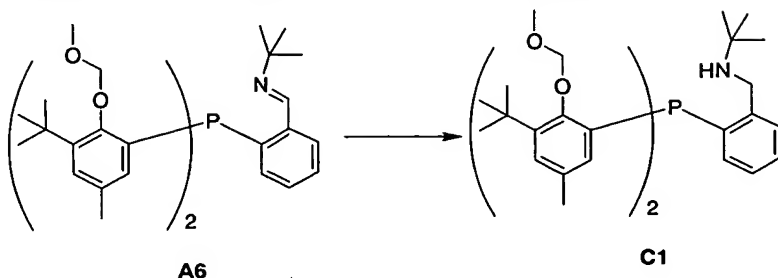
Example B21:

A polymer was produced by the same method as in Example B18 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of methyl aluminoxane. The polymer with a molecular weight (Mw) of 4.00×10^3 , molecular weight distribution (Mw/Mn) of 1.4, a melting point (Tm) of 127.6°C and a number of branches of Me per 1000 carbon atoms of 13 was produced at a rate of 5.10×10^6 g per hour per 1 mol of zirconium by the polymerization reaction.

Example B22:

A polymer was produced by the same method as in Example B18 by polymerization for 11 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of methyl aluminoxane. The polymer with a molecular weight (Mw) of 4.50×10^3 , molecular weight distribution (Mw/Mn) of 1.5, a melting point (Tm) of 129.3°C and a number of branches of Me per 1000 carbon atoms of 20 was produced at a rate of 1.03×10^7 g per hour per 1 mol of zirconium by the polymerization reaction.

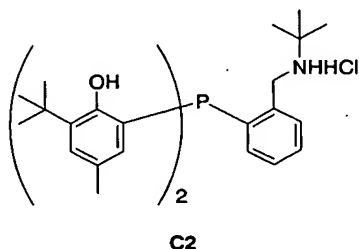
Example C1: Synthesis of compound C1



Sodium borohydride (0.03 g, 0.83 mmol) was added to an ethanol solution (5.76 mL) of compound A6 (0.45 g, 0.75 mmol) at room temperature, and the mixture was stirred for 2 hours. The reaction was stopped by adding deionized water (10.0 mL) and toluene (10.0 mL). After washing the separated organic layer with a saturated aqueous solution (10.0 mL) of sodium chloride, the solvent was removed by evaporation to quantitatively obtain compound C1 as a white solid.

$^1\text{H NMR}(\text{CDCl}_3)$ δ 0.98(9H), 1.32(18H), 2.03(6H), 3.38(6H), 3.3(2H), 5.05(4H), 6.35(2H), 6.78(2H), 7.01-7.20(4H), 7.37(1H)

Example C2: Synthesis of compound C2

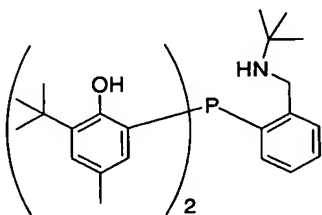


Acetyl chloride (0.32 g, 4.03 mmol) was added to a solution (20.0 mL) of compound C5 (0.49 g, 0.81 mmol) in a mixed solvent (20.0 mL) of ethyl acetate and methanol (1/1) at room temperature, and the mixture was stirred for 15 minutes. Compound C2 was obtained as a white solid (345.0 mg, yield 76.7%) by removing

the solvent by evaporation in vacuum.

^1H NMR(CDCl_3) δ 1.21(9H), 1.40(18H), 2.13(6H), 3.87(2H), 6.38(2H), 7.01-7.37(5H), 7.71(1H), 9.26(2H)

5 Example C3: Synthesis of compound C3

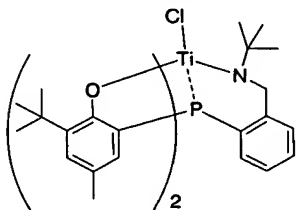


C3

Sodium chloride is added to a tetrahydrofuran solution of compound C6 with stirring. The reaction is stopped by adding deionized solution, the organic layer is separated and compound C3 is obtained by removing the solvent by evaporation.

10

Example C4: Synthesis of complex C4



C4

A tetrahydrofuran solution (3.11 mL) of compound C6 (0.35 g, 0.62 mmol) was added into a tetrahydrofuran solution (2.33 mL) of 60% sodium hydride (0.15 g, 3.72 mmol) at -78°C , and the mixture was warmed to room temperature and then stirred for 1 hour. The reaction mixture was added into a tetrahydrofuran solution (2.33 mL) of titanium tetrachloride bis(tetrahydrofuran) complex (0.21 g, 0.62 mmol) at -78°C . The solution was warmed to room temperature with stirring for 10

hours, and toluene (5.0 mL) was added after removing the solvent by evaporation in vacuum. Complex C4 (258.5 mg, 61.5%) was obtained as a red solid by removing the solvent from the filtrate in vacuum after removing insoluble materials by filtration.

5 $^1\text{H NMR}(\text{C}_6\text{D}_6)$ δ 0.88(9H), 1.34(18H), 1.68(6H), 3.80(2H), 6.60-7.88(8H)
EI-MS 600(M^+)

Examples of polymerization reaction

Example C5:

10 Toluene (5.0 mL) was added to an autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex C4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer was produced
15 at a rate of 3.4×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example C6:

A polymer was produced by the same method as in Example
20 C5, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenyl borane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 1.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example C7:

A polymer was produced by the same method as in Example C5, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and

dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 3.1×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example C8:

A polymer was produced by the same method as in Example C5, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in
10 place of MMAO. The polymer was produced at a rate of 5.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example C9:

15 Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex C4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 20
20 minutes. The polymer with a molecular weight (M_w) of 1.39×10^6 , molecular weight distribution (M_w/M_n) of 8.1 and melting point (T_m) of 122.6°C was produced at a rate of 3.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25 Example C10:

A polymer was produced by the same method as in Example C9, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO.

The polymer with a molecular weight (M_w) of 1.84×10^6 , molecular weight distribution (M_w/M_n) of 19.0 and melting point (T_m) of 124.5°C was produced at a rate of 2.1×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example C11:

A polymer was produced by the same method as in Example C9, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
10 dimethylanilinium terakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 1.94×10^6 , molecular weight distribution (M_w/M_n) of 48.7, melting point (T_m) of 122.9°C and a number of branches of Me per 1000 carbon atoms of 3 was produced at a rate of 2.6×10^6
15 g per hour per 1 mole of titanium by the polymerization reaction.

Example C12:

A polymer was produced by the same method as in Example C9, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
20 triphenylmethyl terakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 3.22×10^6 , molecular weight distribution (M_w/M_n) of 241.9, and melting point (T_m) of 120.6°C was produced at a rate of 4.5×10^6
25 g per hour per 1 mole of titanium by the polymerization reaction.

Example C13:

Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen. After stabilizing at 70°C , ethylene

was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex C4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer with a molecular weight (M_w) of 1.51×10^6 ,
5 molecular weight distribution (M_w/M_n) of 3.4 and melting point (T_m) of 128.0°C was produced at a rate of 2.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example C14:

10 A polymer was produced by the same method as in Example C13, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 3.0×10^5 g per hour per
15 1 mole of titanium by the polymerization reaction.

Example C15:

A polymer was produced by the same method as in Example C13, except that a hexane solution of triisobutyl aluminum (40
20 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium terakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 1.57×10^6 , molecular weight distribution (M_w/M_n) of 4.3, and melting point (T_m) of 121.0°C was produced at a rate
25 of 2.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example C16:

A polymer was produced by the same method as in Example

C13, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl terakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (Mw) of 8.0×10^5 , molecular weight distribution (Mw/Mn) of 2.2, and melting point (T_m) of 120.4°C was produced at a rate of 2.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example C17:

10 Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen. After stabilizing at 130°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex C4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize. The
15 polymer was produced at a rate of 9.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example C18:

A polymer was produced by the same method as in Example
20 C17, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 2.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example C19:

A polymer was produced by the same method as in Example C17, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and

dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 1.20×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example C20:

A polymer was produced by the same method as in Example C17, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
10 triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 7.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

15 Example C21:

Toluene (5.0 mL) and 1-hexene (50 μL) were added to an autoclave under nitrogen. After stabilizing at 70°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmmol) and complex C4 (0.10 μmol) were added to the
20 autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer with a molecular weight (M_w) of 1.5×10^6 , molecular weigh distribution (M_w/M_n) of 3.4, and melting point (T_m) of 118.0°C was produced at a rate of 2.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example C22:

A polymer was produced by the same method as in Example C21, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and

pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 3.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

5 Example C23:

A polymer was produced by the same method as in Example C21, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30
10 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 1.6×10^6 , molecular weight distribution (M_w/M_n) of 4.3, and melting point (T_m) of 121.0°C was produced at a rate of 2.3×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

15

Example C24:

A polymer was produced by the same method as in Example C21, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
20 triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 8.0×10^5 , molecular weight distribution (M_w/M_n) of 2.2 and melting point (T_m) of 120.4°C was produced at a rate of 2.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example C25:

Toluene (5.0 mL) and 1-hexene (40 μL) were added to an autoclave under nitrogen. After stabilizing at 130°C , ethylene was added with compression and was stabilized at 0.60 MPa. MMAO

(100 μmol) and complex C4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 5 minutes. The polymer was produced at a rate of 9.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example C26:

A polymer was produced by the same method as in Example C25, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
10 pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 2.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example C27:

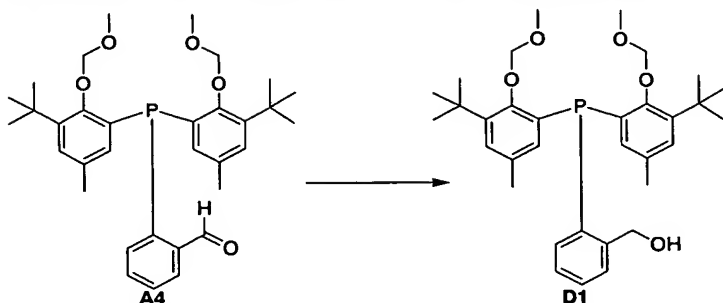
15 A polymer was produced by the same method as in Example C25, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borane (0.30 μmol) were used in place of MMAO. The polymer was produced at
20 a rate of 1.2×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example C28:

A polymer was produced by the same method as in Example
25 C25, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 7.0×10^5 g per hour per 1 mole of titanium by the polymerization

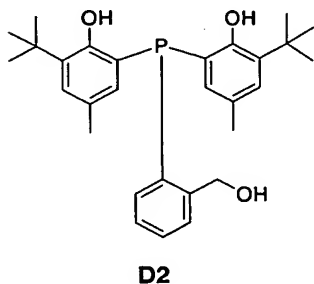
reaction.

Example D1: Synthesis of compound D1



Sodium borohydride (0.95 g, 25.0 mmol) was added to a
5 solution of compound A4 (2.75 g, 5.0 mmol) in ethanol (350 mL)
at room temperature, and the mixture was stirred for 3 hours.
The reaction was stopped by adding deionized water (100.0 mL)
and toluene (100.0 mL), and the organic layer was washed with
a saturated aqueous solution (70 mL) of sodium chloride followed
10 by drying the organic layer over sodium sulfate. A crude product
was obtained as a yellow oil by removing the solvent by evaporation.
The crude product was purified by silica gel column
chromatography (hexane/ethyl acetate = 10/1 → 4/1) to obtain
compound D as a white solid (2.00 g, yield 74.1%).
15 ¹H NMR(CDCl₃) δ 1.39(18H), 2.12(6H), 3.47(6H), 4.80(2H), 5.16-5.21(4H), 6.35(2H),
6.90(1H), 7.14-7.61(5H)

Example D2: Synthesis of compound D2



Acetyl chloride (0.64 g, 8.14 mmol) was added to a solution of compound D1 (1.5 g, 2.71 mmol) in a 1/1 mixed solvent (60.0 mL) of ethyl acetate and methanol at room temperature and the mixture was stirred for 15 hours. Compound D2 was obtained as
5 a white solid (1.06 g, yield 84.5%) by removing the solvent by evaporation in vacuum

^1H NMR(C_6D_6) δ 1.46(18H), 1.82(6H), 5.07(2H), 6.17-7.32(8H), 9.38(2H)

MS Spectrum (EI) 464(M⁺)

10 Example D3: Synthesis of transition metal complex

A 1.57 M hexane solution (1.91 mL) of n-butyl lithium was added dropwise into a tetrahydrofuran solution (7.85 mL) of compound D2 (0.46 g, 1.00 mmol) at -78°C, and the reaction mixture was warmed to room temperature then stirred for 1 hour. The
15 mixed reaction solution was dripped into a tetrahydrofuran solution (7.85 mL) of titanium tetrachloride bis(tetrahydrofuran) complex (0.33 g, 1.00 mmol) at -78°C. The solution was warmed to room temperature and the mixture was stirred for 10 hours. After removing the solvent by evaporation
20 in vacuum, toluene (10.0 mL) was added and insoluble materials were removed by filtration. The transition metal complex was obtained as a red solid (330 mg) by removing the solvent from the filtrate in vacuum.

^{31}P NMR(C_6D_6) δ -1.18

25 MS spectrum (EI) 971

Example of Polymerization Reaction

Example D4:

Toluene (5.0 mL) was added to an autoclave under nitrogen.

After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μ mol) and transition metal complex D3 (0.10 μ mol) obtained in Example D3 were added to the autoclave, and the mixture was allowed to
5 polymerize for 20 minutes. The polymer was produced at a rate of 6.8×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example D5:

10 A polymer was produced by the same method as in Example D4, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenyl borane (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 1.0×10^5 g per hour per
15 1 mole of titanium by the polymerization reaction.

Example D6:

A polymer was produced by the same method as in Example D4, except that a hexane solution of triisobutyl aluminum (40
20 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 3.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example D7:

A polymer was produced by the same method as in Example D4, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and

triphenylmethyl tetrakis(pentafluorophenyl borate ($0.30\ \mu\text{mol}$) were used in place of MMAO. The polymer was produced at a rate of 3.2×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example D8:

Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa.

10 MMAO (100 μmmol) and transition metal complex ($0.10\ \mu\text{mol}$) obtained in Example D3 were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer with a molecular weight (M_w) of 3.1×10^6 , molecular weight distribution (M_w/M_n) of 2.6 and melting point (T_m) of 104.7°C was produced
15 at a rate of 5.7×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example D9:

A polymer was produced by the same method as in Example
20 D8, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane ($0.30\ \mu\text{mol}$) were used in place of MMAO. The polymer was produced at a rate of 1.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example D10:

A polymer was produced by the same method as in Example D8, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and

dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 5.3×10^5 , molecular weight distribution (M_w/M_n) of 10.1 and melting point (T_m) of 135.5°C was produced at a rate of 2.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example D11:

A polymer was produced by the same method as in Example D8, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 1.7×10^6 , molecular weight distribution (M_w/M_n) of 22.8 and melting point (T_m) of 118.0°C was produced at a rate of 2.7×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example D12:

Toluene (5.0 mL) and 1-hexene (50 μL) were added to an autoclave under nitrogen. After stabilizing at 70°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmmol) and transition metal complex (0.10 μmol) obtained in Example D3 were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer with a molecular weight (M_w) of 2.1×10^6 , molecular weight distribution (M_w/M_n) of 5.1 and melting point (T_m) of 115.5°C was produced at a rate of 4.4×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example D13:

A polymer was produced by the same method as in Example D12, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
5 dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 6.9×10^5 , molecular weight distribution (M_w/M_n) of 19.7 and melting point (T_m) of 118.8°C was produced at a rate of 9.0×10^5 g per hour per 1 mole of titanium by the polymerization
10 reaction.

Example D14:

A polymer was produced by the same method as in Example D12, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and
15 triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 8.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

20

Example D15:

Toluene (5.0 mL) and 1-hexene (40 μL) were added to an autoclave under nitrogen. After stabilizing at 130°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa.
25 MMAO (100 μmol) and transition metal complex (0.10 μmol) obtained in Example D3 were added to the autoclave, and the mixture was allowed to polymerize for 5 minutes. The polymer was produced at a rate of 6.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

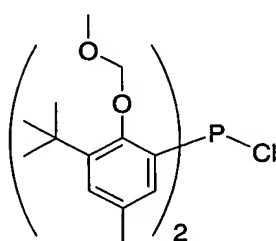
Example D16:

A polymer was produced by the same method as in Example D15, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 3.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

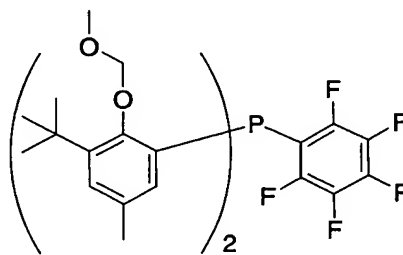
Example D17:

A polymer was produced by the same method as in Example D15, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 6.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example E1: Synthesis of compound E1



A1



E1

A n-butyl lithium solution of (1.56 M) in n-hexane (21.2 mL) was added dropwise into a solution of pentafluorobromobenzene (7.41 g, 30.0 mmol) in diethylether (116.8 mL) at -78°C and the

mixture was stirred for 1 hour. A diethylether solution (50.0 mL) of compound A1 (14.43 g, 30.0 mmol) was added to the mixture, and the resulting mixture was warmed to room temperature with stirring for 5 hours. The reaction was stopped by adding

5 deionized water (100 mL) and toluene (100 mL). After washing the organic layer with a saturated aqueous solution (100 mL) of sodium chloride, the solvent was removed by evaporation to obtain compound E1 as a white solid (17.9 g, yield 98.0%).

^1H NMR(CDCl_3) δ 1.40(18H), 2.18(6H), 3.50(6H), 5.18-5.28(4H),

10 6.53(2H), 7.19(2H)

^{31}P NMR(C_6D_6) δ -30.7

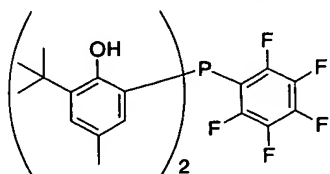
Example E2: Synthesis of compound E1

A 1.56 M hexane solution of n-butyl lithium is added

15 dropwise into a diethyl ether solution of 2-tert-butyl-1-methoxy-4-methylbenzene at -78°C and the mixture was stirred for 1 hour. A diethylether solution of pentafluorophenyl dichlorophosphine is added into the mixed reaction solution, and the solution is warmed to room temperature

20 with stirring. Compound E1 can be obtained by applying the same post-treatment as in Example E1.

Example E3: Synthesis of compound E2



E2

Acetyl chloride (2.65 g, 33.8 mmol) was added to a solution

25 of compound E1 (4.14 g, 6.76 mmol) in a mixed solvent (65.0 mL)

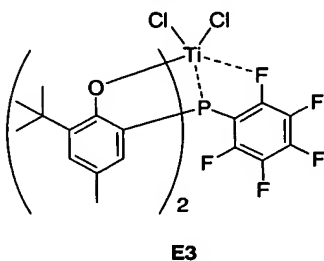
of ethyl acetate/methanol (1/1) at room temperature with stirring for 15 hours. Compound E2 was obtained as a white solid (2.55 g, yield 72.0%) by removing the solvent by evaporation in vacuum.

^1H NMR(CDCl_3) δ 1.40(18H), 2.21(6H), 6.81(2H), 7.17(2H)

5 ^{31}P NMR(C_6D_6) δ -59.6

^{19}F NMR(C_6D_6) δ -161.5, -151.3, -130.7

Example E4: Synthesis of complex E3



A toluene solution of titanium tetrachloride (0.11 g, 0.60 mmol) was dripped into a toluene solution (3.54 mL) of compound E1 (0.31 g, 0.50 mmol), and the solution was warmed to room temperature followed by stirring for 10 hours. Compound E3 was obtained as a red solid (208.7 mg, yield 65.2%) by removing the solvent from the filtrate after filtrating insoluble materials.

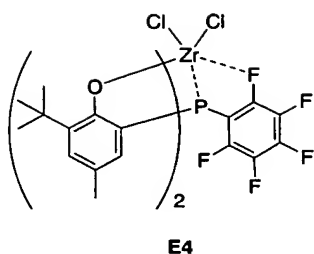
15 ^1H NMR(C_6D_6) δ 1.35-1.44(18H), 1.84-2.01(6H), 6.89-7.01(4H)

^{31}P NMR(C_6D_6) δ 0.36

^{19}F NMR(C_6D_6) δ -161.0, -149.1, -123.1

EI-MS 640(M-1)

20 Example E5: Synthesis of complex E4



A 1.56 M hexane solution (1.03 mL) of n-butyl lithium was added dropwise into a tetrahydrofuran solution (4.73 mL) of compound E2 (0.46 g, 0.80 mmol) at -78°C , and the reaction mixture was warmed to room temperature then stirred for 1 hour. The reaction mixture was added into a tetrahydrofuran solution (10.0 mL) of zirconium tetrachloride bis(tetrahydrofuran) complex (0.30 g, 0.80 mmol). The mixture was warmed to room temperature with stirring for 10 hours. After warming the solution to room temperature with stirring for 10 hours and removing the solvent by evaporation in vacuum, toluene (5.0 mL) was added to the residue. Complex E4 was obtained as a white solid (249.5 mg, yield 45.4%) by removing the solvent from the filtrate after removing insoluble materials by filtration.

^1H NMR(CD_2Cl_2) δ 1.31(18H), 2.21(6H), 6.93(2H), 7.09(2H)

^{31}P NMR(CD_2Cl_2) δ -19.5

^{19}F NMR(CD_2Cl_2) δ -161.6, -151.4, -124.1

EI-MS 684(M $^{+}$)

Example of polymerization reaction

Example E6:

Toluene (5.0 mL) was added to an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and transition metal complex (0.10 μmol) obtained in Example E3 were

added to the autoclave, and the mixture was allowed to polymerize for 3 minutes. The polymer was produced at a rate of 5.99×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

5 Example E7:

A polymer was produced by the same method as in Example E6, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μ mol) were used in place of MMAO.

10 The polymer was produced at a rate of 1.3×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E8:

A polymer was produced by the same method as in Example E6 by polymerization for 5 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium

tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 3.43×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E9:

A polymer was produced by the same method as in Example E6 by polymerization for 3.5 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl

25 tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 4.90×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E10:

Toluene (5.0 mL) and 1-hexene (60 μ L) were added to an autoclave under nitrogen. After stabilizing at 40°C, ethylene
5 was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μ mmol) and complex E3 (0.10 μ mol) were added to the autoclave, and the mixture was allowed to polymerize for 3.6 minutes. The polymer with a molecular weight (M_w) of 7.3×10^4 ,
molecular weigh distribution (M_w/M_n) of 2.7, a melting point
10 (T_m) of 118.2°C and a number of branching of Me per 1,000 atoms of 7 was produced at a rate of 4.52×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E11:

15 A polymer was produced by the same method as in Example E10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μ mol) were used in place of MMAO. The polymer with a molecular weight
20 (M_w) of 2.5×10^4 , molecular weigh distribution (M_w/M_n) of 2.1, a melting point (T_m) of 117.6°C and a number of branching of Me per 1,000 atoms of 4 was produced at a rate of 1.70×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25 Example E12:

A polymer was produced by the same method as in Example E10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl

tetrakis(pentafluorophenyl)borate ($0.30\ \mu\text{mol}$) were used in place of MMAO. The polymer with a molecular weight (M_w) of 3.60×10^4 , molecular weight distribution (M_w/M_n) of 1.8, a melting point (T_m) of 117.2°C and a number of branching of Me per 1,000 atoms of 16 was produced at a rate of 6.02×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E14:

Toluene ($5.0\ \text{mL}$) and 1-hexene ($50\ \mu\text{L}$) were added to an autoclave under nitrogen. After stabilizing at 70°C , ethylene was fed while the ethylene pressure was adjusted at $0.60\ \text{MPa}$. MMAO ($100\ \mu\text{mol}$) and complex E3 ($0.10\ \mu\text{mol}$) were added to the autoclave, and the mixture was allowed to polymerize for 13 minutes. The polymer with a molecular weight (M_w) of 5.6×10^4 , molecular weight distribution (M_w/M_n) of 2.3 and a melting point (T_m) of 127.0°C was produced at a rate of 5.5×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E15:

A polymer was produced by the same method as in Example E14 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum ($40\ \mu\text{L}$, $1.0\ \text{M}$, manufactured by Kanto Chemical Co.) and pentafluorophenylborane ($0.30\ \mu\text{mol}$) were used in place of MMAO. The polymer with a molecular weight (M_w) of 2.8×10^5 , molecular weight distribution (M_w/M_n) of 3.4, a melting point (T_m) of 132.0°C and a number of branching of Me per 1,000 atoms of 1 was produced at a rate of 1.00×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E16:

A polymer was produced by the same method as in Example E14 by polymerization for 6 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 4.9×10^4 , molecular weight distribution (M_w/M_n) of 2.3, a melting point (T_m) of 129.0°C and a number of branching of Me per 1,000 atoms of 6 was produced at a rate of 1.62×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E17:

A polymer was produced by the same method as in Example E14 by polymerization for 6 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 3.60×10^4 , molecular weight distribution (M_w/M_n) of 1.8, a melting point (T_m) of 128.0°C and a number of branching of Me per 1,000 atoms of 8 was produced at a rate of 1.15×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E18:

Toluene (5.0 mL) and 1-hexene (40 μL) were added to an autoclave under nitrogen. After stabilizing at 130°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex E3 (0.10 μmol) were added to the

autoclave, and the mixture was allowed to polymerize for 5 minutes. The polymer was produced at a rate of 8.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

5 Example E19:

A polymer was produced by the same method as in Example E18 by polymerization for 6 minutes, except that a hexane solution of triisobutyl aluminum (4 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.030 μ mol) were used
10 in place of MMAO. The polymer was produced at a rate of 2.00×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example E20:

A polymer was produced by the same method as in Example
15 E18 by polymerization for 6 minutes, except that a hexane solution of triisobutyl aluminum (4 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.030 μ mol) were used in place of MMAO. The polymer was produced at a rate of 8.00×10^5
20 g per hour per 1 mole of titanium by the polymerization reaction.

Example E21:

A polymer was produced by the same method as in Example
E18 by polymerization for 6 minutes, except that a hexane solution
25 of triisobutyl aluminum (4 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.030 μ mol) were used in place of MMAO. The polymer was produced at a rate of 2.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example E22:

Toluene (5.0 mL) was added to an autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene
5 pressure was adjusted at 0.60 MPa. MMAO (100 μ mmol) and complex E4 (0.10 μ mol) were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer was produced at a rate of 2.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

10

Example E23:

A polymer was produced by the same method as in Example E22, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and
15 pentafluorophenylborane (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate of 5.0×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example E24:

20 A polymer was produced by the same method as in Example E22, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced at
25 a rate of 7.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E25:

A polymer was produced by the same method as in Example

E22, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 1.85×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E26:

Toluene (5.0 mL) and 1-hexene (60 μL) were added to an autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex E4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize. The polymer with a molecular weight (M_w) of 3.20×10^5 and molecular weigh distribution (M_w/M_n) of 46.5 was produced at a rate of 1.9×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E27:

A polymer was produced by the same method as in Example E26, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 3.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example E28:

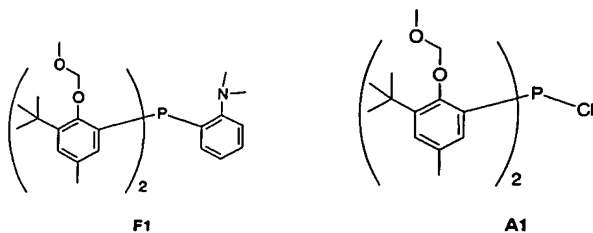
A polymer was produced by the same method as in Example E26, except that a hexane solution of triisobutyl aluminum (40

μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate ($0.30 \mu\text{mol}$) were used in place of MMAO. The polymer with a molecular weight (M_w) of 8.0×10^3 , molecular weight distribution (M_w/M_n) of 1.6 and a number of branches of Me per 1,000 carbon atoms of 42 was produced at a rate of 8.6×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example E29:

A polymer was produced by the same method as in Example E26 by polymerizing for 8 minutes, except that a hexane solution of triisobutyl aluminum ($40 \mu\text{L}$, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate ($0.30 \mu\text{mol}$) were used in place of MMAO. The polymer with a molecular weight (M_w) of 8.6×10^3 , molecular weight distribution (M_w/M_n) of 1.6 and a number of branches of Me per 1,000 carbon atoms of 40 was produced at a rate of 2.51×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example F1: Synthesis of compound F1

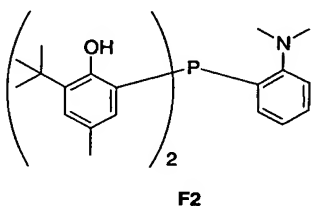


A 1.57 M solution (33.4 mL) of n-butyl lithium was added dropwise into an ether solution (77.3 mL) of 2-N,N-dimethylamino-1-bromobenzene (10.0 g, 50.0 mmol) at -78°C ,

and the mixture was warmed to room temperature then stirred for 1 hour. The solution was cooled to -78°C , and an ether solution (51.0 mL) of compound A1 (24.1 g, 50.0 mmol) was added to the solution followed by warming to room temperature then the resulting mixture was stirred for 3 hours. The reaction was stopped by adding deionized water (100.0 mL) and toluene (100.0 mL). After washing the organic layer with saturated aqueous sodium chloride solution (100 mL), the solvent was removed by evaporation to obtain a desired product as a pale yellow oil. The product was purified by silica gel column chromatography, and compound F1 was obtained as a white solid (13.27 g, yield 46.9%).

^1H NMR(CDCl_3) δ 1.41(18H), 2.10(6H), 2.54(6H), 3.52(6H), 5.25-5.34(4H), 6.29(2H), 6.81-7.30(6H)

Example F2: Synthesis of compound F2

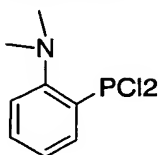


Acetyl chloride (5.89 g, 75.0 mmol) was added to a solution (340 mL) of compound F1 (8.49 g, 15.0 mmol) in a mixed solvent of ethyl acetate/methanol (1/1). Compound F2 was quantitatively obtained as a white solid by removing the solvent by evaporation in vacuum.

^1H NMR(CDCl_3) δ 1.38(18H), 2.10(6H), 3.11(6H), 6.46(2H), 7.19-7.66(6H)

^{31}P NMR(CD_2Cl_2) δ -52.2

Example 3: Synthesis of compound F3



F3

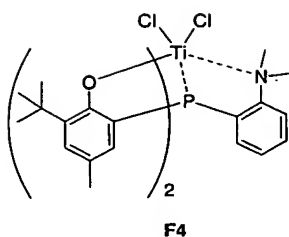
A 1.57 M solution of n-butyl lithium is added dropwise into a tetrahydrofuran solution of

- 5 2-N,N-dimethylamino-1-bromobenzene at -78°C , and the solution was warmed to room temperature and the mixture is stirred for 1 hour. The reaction mixture is dripped into a tetrahydrofuran solution of phosphorous trichloride at -78°C , and the mixture is warmed to room temperature then stirred for 5 hours. Compound
- 10 F3 is obtained by removing the solvent from the filtrate by evaporation in vacuum after removing insoluble materials by filtration.

Example F4: Synthesis of compound F1

- 15 A 1.56 M hexane solution of n-butyl lithium is added dropwise into a tetrahydrofuran solution of 1-methoxy-2-tert-butyl-4-methylbenzne at -78°C , and the reaction mixture is warmed to room temperature then stirred for 1 hour. A tetrahydrofuran solution of compound F3 is added into
- 20 the reaction mixture at -78°C , and the mixture is warmed to room temperature then stirred for 10 hours. Compound F1 is obtained by applying a same post-treatment as in Example F1.

Example F5: Synthesis of complex F4



A toluene solution (8.44 mL) of titanium tetrachloride (0.29 g, 1.55 mmol) was added dropwise into a toluene solution (8.44 mL) of compound F1 (0.73 g, 1.29 mmol) at -78°C , and the mixture was warmed to room temperature then stirred for 10 hours.

5 Complex F4 was obtained as a red brown solid (270 mg, yield 35.1%) by removing the solvent from the filtrate by evaporation in vacuum after removing insoluble materials by filtration.

^{31}P NMR (CD_2Cl_2): δ 26.0

EI-MS: 560 (M-Cl)

10

Example of polymerization reaction

Example F6

Toluene (5.0 mL) was added to an autoclave under nitrogen. After stabilizing at 40°C , ethylene was fed while the ethylene
 15 pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex F4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 8.5 minutes. The polymer was produced at a rate of 1.60×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

20

Example F7:

A polymer was produced by the same method as in Example F6 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto

Chemical Co.) and pentafluorophenylborane ($0.30\ \mu\text{mol}$) were used in place of MMAO. The polymer was produced at a rate of 2.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

5 Example F8:

A polymer was produced by the same method as in Example F6 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum ($40\ \mu\text{L}$, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane ($0.30\ \mu\text{mol}$) were used
10 in place of MMAO. The polymer was produced at a rate of 3.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example F9:

A polymer was produced by the same method as in Example
15 F6 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum ($40\ \mu\text{L}$, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate ($0.30\ \mu\text{mol}$) were used in place of MMAO. The polymer was produced at a rate of 3.0×10^6
20 g per hour per 1 mole of titanium by the polymerization reaction.

Example F10:

Toluene ($5.0\ \text{mL}$) and 1-hexene ($60\ \mu\text{L}$) were added to an autoclave under nitrogen. After stabilizing at 40°C , ethylene
25 was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO ($100\ \mu\text{mol}$) and complex F4 ($0.10\ \mu\text{mol}$) were added to the autoclave, and the mixture was allowed to polymerize for 7 minutes. The polymer with a molecular weight (M_w) of 1.20×10^6 , molecular weight distribution (M_w/M_n) of 99.1 and a number of branches of

Me per 1,000 carbon atoms of 6 was produced at a rate of 1.630×10^7 g per hour per 1 mole of titanium by the polymerization reaction.

Example F11:

5 A polymer was produced by the same method as in Example F10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μ mol) were used in place of MMAO. The polymer was produced at a rate
10 of 2.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Example F12:

 A polymer was produced by the same method as in Example
15 F10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer with a molecular weight (Mw) of
20 1.69×10^6 and molecular weight distribution (Mw/Mn) of 14.0 was produced at a rate of 1.90×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example F13:

25 A polymer was produced by the same method as in Example F10 by polymerization for 20 minutes, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in

place of MMAO. The polymer with a molecular weight (M_w) of 2.18×10^6 and molecular weight distribution (M_w/M_n) of 8.0 was produced at a rate of 1.90×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

5

Example F14:

Toluene (5.0 mL) and 1-hexene (50 μ L) were added to an autoclave under nitrogen. After stabilizing at 70°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μ mol) and complex F4 (0.10 μ mol) were added to the autoclave, and the mixture was allowed to polymerize for 20 minutes. The polymer with a molecular weight (M_w) of 8.1×10^5 , molecular weight distribution (M_w/M_n) of 5.9 and a melting point (T_m) of 116.9°C was produced at a rate of 1.7×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

15

Example F15:

A polymer was produced by the same method as in Example F14 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 9.3×10^5 , molecular weight distribution (M_w/M_n) of 8.9 and melting point (T_m) of 120.6°C was produced at a rate of 1.4×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

25

Example F16:

A polymer was produced by the same method as in Example

F14 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer with a molecular weight (M_w) of 1.0×10^6 , molecular weight distribution (M_w/M_n) of 5.6 and melting point (T_m) of 119.3°C was produced at a rate of 1.4×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

10 Example F17:

Toluene (5.0 mL) and 1-hexene (40 μL) were added to an autoclave under nitrogen. After stabilizing at 130°C , ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. MMAO (100 μmol) and complex F4 (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 5 minutes. The polymer was produced at a rate of 1.1×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example F18:

20 A polymer was produced by the same method as in Example F17 by polymerization, except that a hexane solution of triisobutyl aluminum (4 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of MMAO. The polymer was produced at a rate of 6.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Example F19:

A polymer was produced by the same method as in Example

F17 by polymerization, except that a hexane solution of triisobutyl aluminum (4 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of MMAO. The polymer was produced
5 at a rate of 7.0×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Comparative Example 1:

Toluene (5.0 mL) was added to an autoclave under nitrogen.
10 After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. Methyl aluminoxane (100 μ mmol) and 2,2'-(phenylphosphine)bis(6-tert-butyl-4-methylphenoxy)(tetrahydrofuran)titanium dichloride (0.10 μ mol) were added to the
15 autoclave, and the mixture was allowed to polymerize for 30 minutes. The polymer was produced at a rate of 1.00×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Comparative Example 2:

20 A polymer was produced by the same method as in Comparative Example 1 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and pentafluorophenylborane (0.30 μ mol) were used in place of methyl aluminoxane. The polymer was produced at
25 a rate of 3.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Comparative Example 3:

A polymer was produced by the same method as in Comparative

Example 1 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 1.20×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Comparative Example 4:

10 A polymer was produced by the same method as in Comparative Example 1 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μL , 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μmol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 1.30×10^6 g per hour per 1 mole of titanium by the polymerization reaction.

Comparative Example 5:

20 Toluene (5.0 mL) and 1-hexene (50 μL) were added to an autoclave under nitrogen. After stabilizing at 40°C, ethylene was fed while the ethylene pressure was adjusted at 0.60 MPa. Methyl aluminoxane (100 μmmol) and 2,2'-(phenylphosphine)bis(6-tert-butyl-4-methylphenoxy)(tetrahydrofuran)titanium dichloride (0.10 μmol) were added to the autoclave, and the mixture was allowed to polymerize for 30 minutes. The polymer was produced at a rate of 5.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

Comparative Example 6:

A polymer was produced by the same method as in Comparative Example 5 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and dimethylanilinium tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 6.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

10

Comparative Example 6:

A polymer was produced by the same method as in Comparative Example 5 by polymerization, except that a hexane solution of triisobutyl aluminum (40 μ L, 1.0 M, manufactured by Kanto Chemical Co.) and triphenylmethyl tetrakis(pentafluorophenyl)borate (0.30 μ mol) were used in place of methyl aluminoxane. The polymer was produced at a rate of 7.00×10^5 g per hour per 1 mole of titanium by the polymerization reaction.

20

Industrial Applicability

The transition metal complex having the ligand of the invention is useful as a component of a catalyst for polymerizing olefins. The catalyst has a good polymerization activity and is capable of being used for production of high molecular weight olefin polymers.

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